This is a digital copy of a book that was preserved for generations on library shelves before it was carefully scanned by Google as part of a project to make the world's books discoverable online.

It has survived long enough for the copyright to expire and the book to enter the public domain. A public domain book is one that was never subject to copyright or whose legal copyright term has expired. Whether a book is in the public domain may vary country to country. Public domain books are our gateways to the past, representing a wealth of history, culture and knowledge that's often difficult to discover.

Marks, notations and other marginalia present in the original volume will appear in this file - a reminder of this book's long journey from the publisher to a library and finally to you.

**Usage guidelines**

Google is proud to partner with libraries to digitize public domain materials and make them widely accessible. Public domain books belong to the public and we are merely their custodians. Nevertheless, this work is expensive, so in order to keep providing this resource, we have taken steps to prevent abuse by commercial parties, including placing technical restrictions on automated querying.

We also ask that you:

+ *Make non-commercial use of the files* We designed Google Book Search for use by individuals, and we request that you use these files for personal, non-commercial purposes.

+ *Refrain from automated querying* Do not send automated queries of any sort to Google's system: If you are conducting research on machine translation, optical character recognition or other areas where access to a large amount of text is helpful, please contact us. We encourage the use of public domain materials for these purposes and may be able to help.

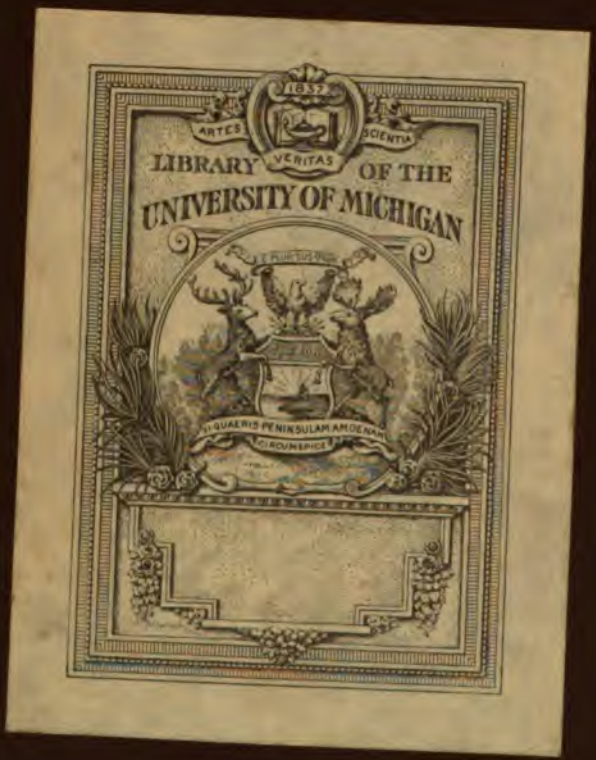
+ *Maintain attribution* The Google "watermark" you see on each file is essential for informing people about this project and helping them find additional materials through Google Book Search. Please do not remove it.

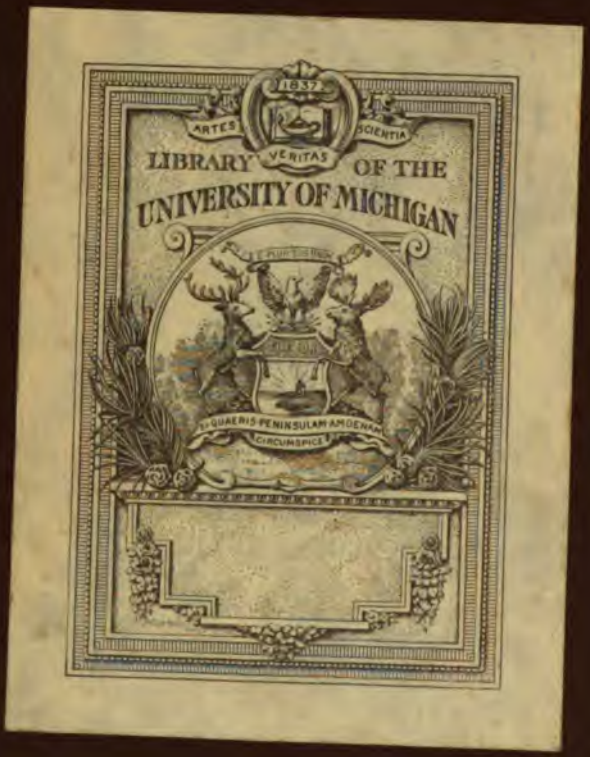
+ *Keep it legal* Whatever your use, remember that you are responsible for ensuring that what you are doing is legal. Do not assume that just because we believe a book is in the public domain for users in the United States, that the work is also in the public domain for users in other countries. Whether a book is still in copyright varies from country to country, and we can't offer guidance on whether any specific use of any specific book is allowed. Please do not assume that a book's appearance in Google Book Search means it can be used in any manner anywhere in the world. Copyright infringement liability can be quite severe.

**About Google Book Search**

Google's mission is to organize the world's information and to make it universally accessible and useful. Google Book Search helps readers discover the world's books while helping authors and publishers reach new audiences. You can search through the full text of this book on the web at `http://books.google.com/`

A 597383 DUPL


1837
ARTES
VERITAS
SCIENTIA
LIBRARY OF THE
UNIVERSITY OF MICHIGAN
SI QUAERIS PENINSULAM AMOENAM
CIRCUMSPICE


1837
ARTES
SCIENTIA
VERITAS
LIBRARY OF THE
UNIVERSITY OF MICHIGAN
SI QUAERIS PENINSULAM AMOENAM
CIRCUMSPICE

# CHEMISTRY:

## GENERAL, MEDICAL, AND PHARMACEUTICAL,

INCLUDING

## THE CHEMISTRY OF THE U. S. PHARMACOPŒIA.

# A MANUAL

ON THE GENERAL PRINCIPLES OF THE SCIENCE, AND THEIR APPLICATIONS IN MEDICINE AND PHARMACY.


BY

JOHN ATTFIELD,

M.A. and PH.D. of the University of Tübingen;

MEMBER OF THE COUNCILS OF THE INSTITUTE OF CHEMISTRY OF GREAT BRITAIN AND IRELAND, AND (LATELY) OF THE CHEMICAL SOCIETY OF LONDON;
PROFESSOR OF PRACTICAL CHEMISTRY TO THE PHARMACEUTICAL SOCIETY OF GREAT BRITAIN;
FORMERLY DEMONSTRATOR OF CHEMISTRY AT ST. BARTHOLOMEW'S HOSPITAL, LONDON;
HONORARY MEMBER OF THE AMERICAN PHARMACEUTICAL ASSOCIATION;
HONORARY MEMBER OF THE COLLEGES OF PHARMACY OF PHILADELPHIA, NEW YORK, MASSACHUSETTS, CHICAGO, AND ONTARIO,
AND OF THE PHARMACEUTICAL ASSOCIATION OF NEW HAMPSHIRE;
HONORARY CORRESPONDING MEMBER OF THE SOCIETY OF PHARMACY OF PARIS;
HONORARY MEMBER OF THE PHARMACEUTICAL SOCIETY OF VICTORIA;
SECRETARY OF THE BRITISH PHARMACEUTICAL CONFERENCE.



*EIGHTH EDITION,*

REVISED BY THE AUTHOR.

PHILADELPHIA:

HENRY C. LEA.

1879.

"But the greatest error of all is, mistaking the ultimate end of knowledge; for some men covet knowledge out of a natural curiosity and inquisitive temper; some to entertain the mind with variety and delight; some for ornament and reputation; some for victory and contention; many for lucre and a livelihood; and but few for employing the Divine gift of reason to the use and benefit of mankind. Thus some appear to seek in knowledge a couch for a searching spirit; others, a walk for a wandering mind; others, a tower of state; others, a fort, or commanding ground; and others, a shop for profit or sale; instead of a storehouse for the glory of the Creator and the endowment of human life."—LORD BACON.

---

Entered according to Act of Congress, in the year 1879, by

HENRY C. LEA,

in the Office of the Librarian of Congress. All rights reserved.

---

COLLINS, PRINTER.

# PREFACE.

THE short title on the back of a book, and even the words on the title-page, are generally, and even necessarily, imperfect descriptions of the contents, and hence not unfrequently induce at the outset misconceptions in the minds of readers. The Author of "Chemistry, General, Medical, and Pharmaceutical" would at once state, therefore, that his sole aim is to teach the general truths of chemistry to medical and pharmaceutical pupils. So far as laws and principles are concerned, the book is a work on General Chemistry; but, inasmuch as those laws and principles are elucidated and illustrated by that large portion of chemistry which is directly interesting to medical practitioners and pharmacists, the book may be said to be a work on Medical Chemistry and Pharmaceutical Chemistry. Only in this conventional sense would the author speak of Medical and Pharmaceutical Chemistry; for the truths of Chemistry are the same for all students—crystalline verities which cannot be expanded or compressed to suit any class of workers. The leading principles of the science, however, can as easily be illustrated by or deduced from those facts which have interest as from those which have little or no special interest to the followers of medicine and pharmacy. The grand and simple leading truths of chemistry, the lesser truths or principles, and nearly all the interesting relationships of elements and compounds, in a word, the *science* of chemistry, can be taught to medical and pharmaceutical students with little other aid than that afforded by the materials which lie in rich abundance all around these workers. Such a mode of teaching "the general principles of the science, and their applications in medicine and

pharmacy," is adopted in this volume. It is a mode which greatly increases the usefulness of the science to the class addressed, while it in no way diminishes the value of chemistry as an instrument of mental culture—an instrument which sharpens and expands the powers of observation, which enlarges and strengthens memory and imagination, which gives point to the perceptive faculties, and which develops and elaborates the powers of thought and of reason.

This manual is intended, then, as a systematic exponent of the general truths of Chemistry, but is written solely for the pupils, assistants, and principals engaged in medicine and pharmacy. It will be found equally useful as a reading-book for gentlemen having no opportunities of attending lectures or performing experiments, or, on the other hand, as a text-book for college pupils; while its comprehensive Index, containing six thousand references, will fit the work for after-consultation in the course of business or professional practice.

From other chemical text-books it differs in three particulars: first, in the exclusion of matter relating to compounds which at present are only of interest to the scientific chemist; secondly, in containing more or less of the chemistry of every substance recognized officially, or in general practice, as a remedial agent; thirdly, in the paragraphs being so cast that the volume may be used as a guide in studying the science experimentally.

The order of subjects is that which, in the author's opinion, best meets the requirements of medical and pharmaceutical students in Great Britain, Ireland, America, and the English Colonies. Introductory pages are devoted to a few leading properties of the elements. A review of the facts thus unfolded affords opportunity for stating the views of philosophers respecting the manner in which these elements influence each other as components of terrestrial matter. The consideration in detail of the relations of the elementary and compound radicals follows, synthetical and analytical bearings being pointed out, and attention frequently directed to connecting or underlying truths or general principles. The chemistry of substances naturally

associated in vegetables and animals is next considered. Practical toxicology and the chemical as well as microscopical characters of morbid urine, urinary sediments, and calculi are then given. The concluding sections form a laboratory-guide to the chemical and physical study of quantitative analysis. In the appendix is a long table of tests for impurities in medicinal preparations—also a short one of the saturating-powers of acids and alkalies, designed for use in prescribing and dispensing.

In the course of the treatment outlined in the preceding paragraph, it will be observed that the whole of the elements are first noticed very shortly, to give the pupil a general view of his course of study, and afterwards at length and thoroughly; that the chemistry of the common metallic radicals precedes that of the rarer, and that the sections on the acidulous radicals are similarly divided; while the basylous radicals are arranged according to analytical relations, the common acidulous according to exchangeable value or quantivalence, and rarer acidulous radicals alphabetically. By this plan the more important facts and principles are repeatedly brought under consideration, the points of view, however, differing according as interest is concentrated on physical, synthetical, analytical, or quantitative properties. This arrangement of matter was adopted, also, partly from the belief that the separate and general truths of chemistry never enter the mind in the order of any scientific classification at present possible. In the current state of chemical knowledge, consistency in the methodical arrangement even of elements can only be carried out in one direction, and is necessarily accompanied by inconsistencies in other directions, a result most perplexing to learners, and hence totally subversive of the chief advantage of classification. For this reason the writer has preferred to lead up to, rather than follow, scientific classification—has allowed analogies and affinities to suggest, rather than be suggested by, classification. Among the acidulous radicals, especially, any known system of classification would have given undue prominence to one set of relations and undeserved obscurity to others. Then, by separating more

important from less important matter, instruction is adapted to the wants of gentlemen whose opportunities of studying chemistry vary greatly, and are unavoidably insufficient to enable them to gain a knowledge of the detail of the science. One great advantage of the mode of treatment is that difficulties of nomenclature, notation, chemical constitution, and even those arising from conventionality of language, are explained as they arise, instead of being massed under the head of "Introductory Chapters," "Preliminary Considerations," or "General Remarks," which are not unfrequently too difficult to be understood by a beginner, too voluminous to be remembered except by the aid of subsequent lessons, and are consequently the cause of much trouble and confusion. This plan has also admitted of greater prominence being given to "The General Principles of Chemical Philosophy," the only section to which the student is asked frequently to return until he finds himself naturally employing those principles in the interpretation of the phenomena obtained by experiment. An elementary knowledge of the subjects of Gravitation, Heat, Light, Sound, Electricity, and Magnetism cannot be too strongly recommended to the student of Chemistry. The first portion of this Manual would have been devoted to an exposition of these branches of Physics, so far as they bear on Chemistry, did not the many special books on Physics render such a course unnecessary. Quantitative chemical analysis frequently involving determinations of temperature, specific gravity, and atmospheric pressure, a few paragraphs on these subjects are made introductory to the sections on quantitative operations.

The theories that matter consists of molecules and that molecules consist of atoms are freely adopted in this book, the author believing that, in the present state of knowledge and education, philosophic conceptions regarding chemistry can only be taught to medical, pharmaceutical, and the great majority of general students by some objective aid.

The chemical notation of the work is in accordance with modern theories. Equations illustrative of pharmacopœial processes have a name attached to each formula.

Chemical nomenclature has been modernized to the extent of defining the alkali-metal salts and the earthy compounds as those of potassium, sodium, ammonium, barium, calcium, magnesium, and aluminium, instead of potash, soda, ammonia, baryta, lime, magnesia, and alumina. The author confidently believes that this change, founded on views now adopted by all prominent writers on chemistry, and used in the Pharmacopœia of the United States, will be accepted and become popular in medicine and pharmacy; it is a step in the direction of simplicity and consistency, and involves far less hypothesis than is contained in the old system. The name nitrate of potash, for example, was based on the pure assumption that nitre contained oxide of potassium or potash and nitric anhydride, then erroneously termed nitric acid. By the modern name nitrate of potassium, all that is intended to be conveyed is that nitre contains the element common to all potassium compounds, and the group of elements common to all nitrates. Under the old method, students always experienced difficulty in distinguishing salts of the metal from salts of its oxide—salts of potassium, for instance, from salts of potash; under the new view no such difficulty arises. Names such as potassium nitrate or potassic nitrate are also consistent with modern views, but for general adoption are too unlike the original. The contractions in Latin for names like "nitrate of potassium" are identical with those names resembling "nitrate of potash"; an accidental circumstance that will much facilitate the general introduction of the former among medical practitioners and pharmacists, and a practical advantage that must determine the choice over the other chemically equivalent names just mentioned. The author ventures to express some gratification that his use and advocacy of this system since the first edition of this Manual was published in 1867 has resulted in its adoption, in 1873, in the "Pharmacopœia of the United States," and in the recommendation, from all medical, chemical, and pharmaceutical authorities in Great Britain, of its adoption in the next "British Pharmacopœia." Pharmacy in these two countries will thus sooner or later, in the important matter of chemical nomenclature, be in accord with the current state of chemical science.

The Metric System of Weights and Measures (that which, doubtless, is destined to supersede all others) is alone used in the sections on Quantitative Analysis. In other parts of the Manual avoirdupois weights and imperial measures are employed.

It is hoped that the numerous etymological references scattered throughout the following pages will be found useful. Words in Greek have been rendered in English characters, letter for letter. The word "official" is used throughout for things recognized officially by the compilers of Pharmacopœias; "officinal" in its original application to the officina or shop.

Students are strongly recommended to test their progress by frequent examination. To this end appropriate questions are appended to each subject.

The Author's ideal of a manual of Chemistry for medical and pharmaceutical students is one in which not only the *science* of chemistry is taught, but in which the chemistry of every substance having interest for the followers of medicine and pharmacy is noticed at more or less length in proportion to its importance, and at least its position in relation to the leading principles of chemistry set forth with all attainable exactness. The extent to which he has realized this ideal he leaves to others to decide. Such a work will doubtless in certain parts partake of the character of a dictionary; but this is by no means a fault, especially if a good index be appended; for the points of contact between pure and applied chemistry are thus multiplied, and abundant outlets supplied, by which a lover of the science may pass into other chemical domains by aid of other guides, or even into the regions of original research. Among the rarer alkaloids, bitter bodies, glucosides, salts or organic radicals, solid fats, fixed oils, volatile oils, resins, oleo-resins, gum-resins, balsams, and coloring-matters, mentioned in this volume, will be found many such points whence the ardent student may start for the well-known obscure or untrodden paths of scientific chemistry.

Within twelve years a demand has arisen for eight large editions of this Manual. The First, in 1867, was intended as a handbook of practical chemistry only; but the notes and remarks made respecting most of the experiments were found to be so useful by students that this portion of the volume was in the Second Edition (1869) sufficiently extended to render the book more fairly complete in itself. In response to a call from professional friends in the United States in 1870, the work was revised by the Author for the followers of medicine and pharmacy in America, the chemistry of the Preparations and Materia Medica of the United States Pharmacopœia being introduced, and such other adaptations included as to form a Third Edition. A Fourth was presented to English workers in the autumn of 1872, and founded on the Fourth, a Fifth Edition for American students in 1873. A very large Sixth Edition was published in England in 1875, and a Seventh in America in 1876.

The present, Eighth, edition contains such alterations and additions as seemed necessary for the demonstration of the latest developments of chemical principles and the latest applications of chemistry in Pharmacy. The Author has bestowed assiduous labor on the revision, and the extent of the information thus introduced may be estimated from the fact that the Index contains three hundred new references relating to the additional material. The work now includes the whole of the chemistry of the United States Pharmacopœia, of the British Pharmacopœia, and of the Pharmacopœia of India.

Engravings, by G. Pearson, Esq., of most of the important pieces of apparatus employed in studying chemistry experimentally, are reproduced in this Eighth Edition. The twenty microscopic views are after drawings by Tuffen West, Esq.

17 Bloomsbury Square, London,
January 1st, 1879.

## ADVICE TO STUDENTS.

It is unnecessary to advise you to avoid studying merely by way of "preparation for examination." You will not so mistake the means for the end. You are studying to fit yourself for your position in the world. Work diligently, study thoughtfully and deliberately, above all be thorough; otherwise your knowledge will be transient, and will be unaccompanied by that enlightenment of the understanding, that mental training, mental discipline, and general elevation of the intellect which constitute, in a word, education. When you are thus educated, you will with ease and pleasure pass any examination in the knowledge you have thus acquired.

All authorities on education, whether statesmen, teachers, or examiners, regard "Examination," even by the most highly skilled "Board," with ample time at its disposal and a wide area from which to select questions, as but a partial test of knowledge, and an extremely imperfect test of education. It is the best, however, that has been devised, and is especially useful when, following instead of leading education, it is restricted to the subjects of a well-defined, earnestly followed, compulsory curriculum of study.

Students! as men of honor, take care that any inefficiencies inseparable from "examination," are abundantly compensated by the extent and precision of your knowledge, and by the soundness and thoroughness of your whole education.

---

## APPARATUS FOR EXPERIMENTS IN ANALYSIS.

List of Apparatus suitable for a three months' course of practical chemistry in the summer session of medical schools, or for any similar series of lessons—including the preparation of elementary gases, analytical reactions of common metals and acidulous radicals, analysis of single salts, chemical toxicology, and the examination of urine, urinary sediments, and calculi:—

One dozen test-tubes.
Test-tube stand.
Test-tube cleaning-brush.
A few pieces of glass tubing, 8 to 16 in. long, with a few inches of India-rubber tubing to fit.
Small flask.
Two small beakers.
Two small funnels.
Two watch-glasses.
Two or three glass rods.
Wash-bottle.
Small pestle and mortar.
A 2-pint earthenware basin.
A 2-inch and a 3-inch evap. basin.
Two porcelain crucibles.
Blowpipe.
Crucible tongs.
Round file.
Triangular file.
Small retort-stand.
Sand-tray.
Wire triangles.
Platinum wire and foil.
Test-paper.
Filter-paper.
Towel.
Two dozen corks.

(*This set, packed in a deal-box, can be obtained of any chemical-apparatus maker for about seven dollars.*)

### APPARATUS FOR EXPERIMENTS IN SYNTHESIS AND ANALYSIS.

A larger set, suitable for the performance of most of the synthetical as well as analytical experiments described in this manual:—

- A set of evaporating-basins, of the following sizes:—
  - One 8½-inch.
  - One 7¼-inch.
  - One 6½-inch.
  - One 4-inch.
  - Two 3-inch.
- One retort-stand and three rings.
- Two test-glasses.
- One half-pint flask.
- One half-quire filter-paper.
- Two porcelain crucibles.
- One measure-glass, 5 oz.
- Blowpipe, 8-inch, Black's.
- Two glass funnels.
- One doz. test tubes (German glass)
- One test-tube brush.
- One pair of 8-inch brass crucible-tongs.
- Two soup-plates.
- One flat plate.
- Two spatula knives.
- One pair of scissors.
- One round file.
- One triangular file.
- One half-pound glass rod.
- One half-pound glass tubing.
- One ft. small India-rubber tubing.
- Three dozen corks of various sizes.
- Platinum wire and foil.
- Test-papers.
- A nest of three beakers.

(*This set, packed in a case, can be obtained of any chemical-apparatus maker for about twelve dollars.*)

A sponge, towels, and note-book may be included.

### FURNITURE OF A LABORATORY.

The following apparatus should be ready to the hands of students following an extended course of practical chemistry, in a room set apart for the purpose:—

- A bench or table and stool.
- Water-supply and waste-pipe.
- A cupboard attached to a chimney with an outward draught.
- A furnace fed with coke; tongs, hot-plate, or sand-bath, etc.
- A waste box.
- Shelves for chemicals and other materials in jars or bottles.
- Gas-supply and lamp with flexible tube (or a spirit lamp and spirit).
- Test-tube rack, two dozen holes.
- Iron stand or cylinder for supporting large dishes.
- Iron adapters for fitting dishes to cylinder.
- Pestle and mortar, 5 or 6 inches.
- One 6-inch funnel.
- Brown pan, 1- or 2-gallon.
- White jug, 1-gallon.
- Water-bottle, quart.
- Twenty-eight test-bottles, 6-oz.

Other articles, such as flasks, retorts, receivers, condensers, large evaporating-dishes, may be obtained as wanted. In Quantitative Analysis the apparatus described in the sections on that subject will be required.

## REAGENTS.

Certain chemicals are used so frequently in analytical processes that it is desirable to have small quantities placed in bottles in front of the operator. As these reagents or "tests" are generally employed in a state of solution, nearly all the solid salts may at once be dissolved (in distilled water). The bottles employed should be well stoppered, and of 5 or 6 ounces capacity. The bottles should not be more than three-quarters full; single drops, if required, can then be poured out with ease and precision. The following list of test solutions is recommended; directions for methods of preparing those not readily purchasable will be found by referring to the Index:—

| | |
|---|---|
| Sulphuric Acid, strong. | Sol. of Potash, 5 per cent. or B. P. |
| Nitric Acid, strong. | " Soda, 5 to 15 per cent. |
| Hydrochloric Acid, strong. | " Ammon., 10 per ct. or B.P. |
| Acetic Acid, strong. | Lime-water, saturated. |

The next nine may contain about 10 per cent. of solid salt:—

| | |
|---|---|
| Carbonate of Ammonium, with a little solution of Ammonia added. | Sulphydrate of Ammonium. |
| | Chloride of Barium. |
| Chloride of Ammonium. | Chloride of Calcium. |
| Phosphate or Arseniate of Ammonium. | Phosphate of Sodium. |
| | Neutral Chromate. |

The succeeding seven may have a strength of about 5 per cent.:—

| | |
|---|---|
| Ferrocyanide of Potassium. | Perchloride of Iron. |
| Ferridcyanide of Potassium. | Nitrate of Silver. |
| Iodide of Potassium. | Perchloride of Platinum. |
| Oxalate of Ammonium. | |

---

## LIST OF CHEMICALS.

List of chemicals necessary for the practical study of the non-metallic elements mentioned on pp. 13 to 31. The quantities are sufficient for several experiments.

| | | | |
|---|---|---|---|
| Chlorate of Potassium | 1 oz. | Phosphorus | ½ oz. |
| Black Oxide of Manganese | 1 oz. | Hydrochloric Acid | 1 oz. |
| Zinc | 1 oz. | Sulphur | ½ oz. |
| Oil of Vitriol | 2 oz. | Iodine | ¼ oz. |

List of chemicals necessary for the *analytical* study of the metallic and acidulous radicals (pages 61 to 378). The quantities will depend on the frequency with which experiments are repeated or analysis performed; those mentioned are sufficient for one or two

students. The articles are given in the order in which they will be required. The eight substances mentioned in the above list are included.

| | | | |
|---|---|---|---|
| The set of test solutions described on the previous page. | | Nitrate of Strontium | ¼ oz. |
| Carbonate of Potassium | 1 oz. | Black Oxide of Manganese | ½ lb. |
| Tartaric Acid | 1 oz. | Chloride of Manganese | ¼ oz. |
| Litmus | ¼ oz. | Chloride of Cobalt | 50 grs. |
| Sulphate of Magnesium | 1 oz. | Nitrate of Nickel | ¼ oz. |
| Sulphate of Zinc | 1 oz. | Chloride of Chromium | ¼ oz. |
| Alum | 1 oz. | Gold leaves | 2 or 3 |
| Sulphide of Iron | 1 lb. | Chloride of Cadmium | ¼ oz. |
| Oak-galls | 1 oz. | Nitrate of Bismuth | ¼ oz. |
| Sulphocyanate of Potassium | ¼ oz. | Bromide of Potassium | ¼ oz. |
| White Arsenic | ½ oz. | Starch | 1 oz. |
| Zinc | ½ lb. | Nitrate of Potassium | 1 oz. |
| Charcoal | ½ lb. | Copper borings or turnings | 1 oz. |
| Sulphate of Iron | 1 oz. | Indigo | ¼ oz. |
| Copper Foil | 1 oz. | Chlorate of Potassium | 1 oz. |
| Sulphate of Copper | 1 oz. | Iodine | ½ oz. |
| Tartar Emetic | ½ oz. | Spirit of Wine | 1 oz. |
| Mercury | 1 oz. | Sulphur | 1 oz. |
| Corrosive Sublimate | ½ oz. | Acid Oxalate of Potassium | 1 oz. |
| Calomel | ½ oz. | Citric Acid | 1 oz. |
| Tin | 1 oz. | Phosphorus | 1 oz. |
| Bicarbonate | 1 oz. | Borax | 1 oz. |
| Acetate of Lead | 1 oz. | Turmeric | 1 oz. |
| Cyanide of Potassium | ½ oz. | Benzoic Acid | 50 grs. |
| Hyposulphite of Sodium | 1 oz. | Fluor Spar | 1 oz. |
| A Lithium Salt | 10 grs. | Tannic Acid | 50 grs. |
| | | Gallic Acid | 50 grs. |
| | | Pyrogallic Acid | 50 grs. |

The quantities of materials required for the study of chemistry *synthetically* will necessarily vary with the desires and tastes of the operator, or according to the number and requirements of students working together.

# CONTENTS.


PAGE

PREFACE . . . . . iii

ADVICE TO STUDENTS . . . . . xi

APPARATUS . . . . . xi

REAGENTS . . . . . xiii

LISTS OF CHEMICALS . . . . . xiii

INTRODUCTION . . . . . 13

GENERAL PROPERTIES OF THE NON-METALLIC ELEMENTS . . . . . 15

SYMBOLS AND DERIVATION OF NAMES OF ELEMENTS . 31

THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY 36

COMMON METALLIC ELEMENTS, THEIR OFFICIAL PREPARATIONS, AND TESTS :—

SALTS OF POTASSIUM, SODIUM, AMMONIUM, BARIUM, CALCIUM, MAGNESIUM, ZINC, ALUMINIUM, IRON, ARSENICUM, ANTIMONY, COPPER, MERCURY, LEAD, SILVER . . . . . 60

ANALYTICAL CHARTS, FOR ORDINARY METALS . . . 220

RARER METALLIC ELEMENTS, THEIR OFFICIAL PREPARATIONS, AND TESTS :—

SALTS OF LITHIUM, STRONTIUM, MANGANESE, COBALT, NICKEL, CHROMIUM, TIN, GOLD, PLATINUM, CADMIUM, BISMUTH . . . . . 226

ANALYTICAL CHARTS, FOR ALL METALS . . . . . 258

COMMON ACIDULOUS RADICLLS, OFFICIAL ACIDS, AND TESTS :—

CHLORIDES, BROMIDES, IODIDES, CYANIDES, NITRATES, CHLORATES, ACETATES, SULPHIDES, SULPHITES, SULPHATES, CARBONATES, OXALATES, TARTRATES, CITRATES, PHOSPHATES, BORATES . 263

PAGE

Salts of Rarer Acidulous Radicals :—

Benzoates, Cyanates, Formates, Hippurates, Ferrocyanides, Ferridcyanides, Fluorides, Hypophosphites, Hyposulphites, Lactates, Malates, Meconates, Metaphosphates, Nitrites, Phosphites, Pyrophosphates, Silicates, Sulphocyanates, Tannates, Gallates, Urates, Valerianates . . . . . 335

Analytical Chart for Acidulous Radicals . . . 366

Systematic Analysis . . . . . 369

Alkaloids, Amylaceous and Saccharine Substances, Glucosides, Alcohol, and Allied Bodies, Albumenoid and Gelatigenous Substances, Pepsine, Fatty Bodies, Resinoid Substances, Coloring-matters . . . . . 378

Toxicology . . . . . 483

Examination of Morbid Urine and Calculi . . . 493

Official Galenical Preparations . . . . . 507

Official Chemical Preparations . . . . . 509

Quantitative Analysis :—

Introductory Remarks . . . . . 510

Measurement of Temperature . . . . . 513

Estimation of Weight . . . . . 518

Weights and Measures . . . . . 519

Specific Gravity . . . . . 529

Correction of the Volume of Gases for Pressure and Temperature . . . . . 536

Volumetric Analysis . . . . . 543

Gravimetric Analysis . . . . . 567

Dialysis . . . . . 610

Appendix :

Table of Official Tests for Impurities in Preparations of the British Pharmacopœia . 615

Saturation Tables . . . . . 623

The Elements, their Symbols and Atomic weights . . . . . 635

Index . . . . . 637

# CHEMISTRY:

## GENERAL, MEDICAL, AND PHARMACEUTICAL.

---

### INTRODUCTION.*

THE infinite variety of solid, liquid, and gaseous substances of which our earth and atmosphere are composed, may be resolved with more or less difficulty into distinct forms of matter appropriately termed Elements, for by no known means can they be further decomposed. Sixty-seven elements have been proved to exist. A few (such as gold) occur naturally in the uncombined state, but the greater number are combined in so subtle a manner as to conceal them from ordinary methods of observation. Thus none of the common properties of water indicate that it is composed of two elements, both gases, but differing much from each other: nor can the senses of sight, touch, and taste, or other common means of examination, detect in their concealment the three elements of which sugar is composed. The art by which these and all other compound substances are resolved into their elements is termed Chemistry, a name derived possibly from the Arabic word *kamai*, to conceal.† The *art* of chemistry also includes the construction of compounds from elements, and the conversion of substances of one character into those of another. The general principles

---

* Students using this book as a guide in following chemistry practically should read the first three pages, and then commence work by preparing oxygen. All students should read the Preface.

† The idea that common metals contained valuable metals concealed within them was the one seed from which mainly sprung chemical knowledge. The men who endeavored to find the secret of such concealment were appropriately termed *alchemists*, and their efforts spoken of as *alchemy* (*al kimia*, from *kamai*, to conceal). Their persistent labors, generation after generation, were unsuccessful so far as the transmutation of baser metals into gold was concerned, yet were invaluable to posterity. For new substances were discovered and truths of nature unveiled; from these discoveries multiplication of discoveries resulted, and thus grew the still-growing branch of knowledge called Chemistry.

or leading truths relating to the elements, to the manner in which they severally combine, and to the properties of the compound substances formed by their union, constitute the *science* of chemistry.*

From these few words concerning the nature of the art and science of chemistry, it will be seen that in most of the occupations that engage the attention of man chemistry plays an important part—in few more so than in the practice of Therapeutics† and Pharmacy.‡

---

* Persons who practise the art and science of Chemistry are known as Chemists. Some two hundred or more years ago, and before chemistry was a science, the "chemists" were the makers or vendors of chemicals, then only used as medicines. They were the successors of the Alchemists. In Great Britain these chemists and the herbalists, otherwise drug-grocers, otherwise druggists, gradually associated to form the "Chemists and Druggists." Between the "Chemist and Druggist" and the Physician there existed the Apothecary—the putter together of medicines or compounder of physicians' prescriptions. The Apothecary has since become a medical practitioner, prescriptions now being "made up" by the Chemist and Druggist. The latter in Great Britain, since the year 1868, is allowed the *title* of *Chemist and Druggist*, his higher title being *Pharmaceutical Chemist;* these respective designations he legally assumes on passing the Minor and Major Examinations, conducted by the Pharmaceutical Society of Great Britain in accordance with the provisions of the Pharmacy Acts of 1852 and 1868. The whole class is often spoken of as that of *Pharmacists* or *Pharmaceutists*, terms also used in the United States. Other classes of chemists are the Analytical Chemists, who give special attention to Analysis; Manufacturing Chemists, who restrict their labors to the preparation of chemicals; while others devote a portion of their knowledge and energies to Chemical Education or to Chemical Research, or are appealed to as Consulting Chemists by the persons, firms, corporations, or governments needing chemical advice respecting industrial processes, hygienic matters, etc. The callings of the Consulting and Analytical Chemist are generally united, and the professional gentlemen who follow these conjoint avocations also not unfrequently occupy professorial or other tutorial positions, sometimes adding to these labors more or less work at original chemical research. In England, Scotland, and Ireland, nearly all the leading professional chemists are *Fellows of the Institute of Chemistry of Great Britain and Ireland.*

† Therapeutics (θεραπευτικος, *therapeutikos*, from θέραπευω, *therapeuo*, to nurse, serve, or cure) is that branch of medicine which treats of the application of remedies for diseases; it includes dietetics. The therapeutist also takes cognizance of hygiene, that department of medicine which respects the preservation of health.

‡ Pharmacy (from φάρμακον, *pharmakon*, a drug) is the generic name for the operations of preparing or compounding medicines, whether performed by the Medical Practitioner or by the Chemist and Druggist. It is also sometimes applied, like the corresponding term "Surgery," to the apartment in which the operations are conducted.

Air, water, food, drugs, and chemicals, in short all material substances, are composed of a few elements. An intimate knowledge of the properties of these, and of the various substances they form by combining with each other, a knowledge of the power or force (the chemical force, or chemical affinity) by which the elements contained in those compounds are held together, and an application of such knowledge to Pharmacy and Medicine, must be the objects sought to be attained by the learner, for whom this work has been especially written.

---

*The Elements.*—Of the sixty-seven known elements thirty-nine are of medical or pharmaceutical interest; of these, about two-thirds are metals, and one-third non-metals; the remainder are so seldom met with in nature as to have received no practical application either in medicine, art, or manufacture. Before intimately studying the elements, it is desirable to acquire some general notions concerning them: such a procedure will also serve to introduce the practical student to his apparatus, and make him better acquainted with the various methods of manipulation.*

*Metallic Elements.*—With regard to the metallic elements, it may be safely assumed that the reader has sufficient knowledge for present purposes; but little, therefore, need now be said respecting them. He has an idea of the appearance, relative weight, hardness, etc., of such metals as gold, silver, copper, lead, tin, zinc and iron. If he has not a similar knowledge of mercury, antimony, arsenicum, platinum, nickel, aluminium, magnesium, potassium, and sodium, he should commence his studies by seeing and handling specimens of each of these metals.

*Non-Metallic Elements.*†—With regard to the non-metallic

---

* This allusion to apparatus need not discourage the youngest pupil. With the aid of a few phials, wine-glasses, or other similar vessels always at hand, he may, by studying the following pages, learn the chemical reactions which are constantly occurring in the course of making up medicines, understand the process by which medicinal preparations are manufactured, and detect adulterations, impurities, or faults of manufacture. Among the substances used in medicine, will be found nearly all the chemicals required. If, in addition, a dozen test-tubes and a few feet of glass tubing be procured, many of the experiments described may be performed. For full lists of apparatus and chemicals see introductory pages.

† These bodies are sometimes termed *metalloids* (from μέταλλον, *metallon*, a metal, and εἶδος, *eidos*, likeness); but the name is not appropriate, for the non-metallic elements have no likeness to metals.

elements, it is here supposed that the student has no general knowledge. He should commence his studies therefore by a series of operations as follows, on eight out of their number.

## OXYGEN.

*Preparation.*—As oxygen is the most abundant element in nature, forming, though in a combined state, about one-half of the whole weight of our globe, it may safely be assumed that this element can readily be obtained in the free condition in a state of purity. In fact, the air itself contains about one-fifth of its bulk of oxygen, though from the air it cannot be separated sufficiently easily and readily for experimental purposes. It is preferable to apply heat—that force which will often be noticed as antagonistic, so to speak, to chemical union; heat generally separating particles of matter further from each other, while chemical attraction tends to bind them closer together—it is better to heat certain compounds containing oxygen; the latter is then evolved in its normal, natural condition of gas. Several substances, when heated, yield oxygen; but, for convenience and economy, the crystalline body known as chlorate of potassium is best fitted for the experiment. The size and form of the vessel in which to heat it will mainly depend on the quantity required; but for the purposes of the student the best is a *test-tube*, an instrument in constant requisition in studying practical chemistry. It is simply a thin tube of glass, a few inches in length, and half or three-quarters of an inch in diameter, closed by fusion at one end. It is made of thin glass, in order that it may be rapidly heated or cooled without risk of fracture. (See Fig. 1.)

*Outline of the Process.*—Heat chlorate of potassium (say, as much as will lie on a shilling) in a test-tube, by means of a spirit- or gas-flame; gaseous oxygen is quickly evolved. Before applying heat, however, provision should be made for collecting the gas. (See Fig. 3.)

*Collection of Gases.*—Procure a piece of glass tubing about the thickness of a quill pen, and a foot or eighteen inches long, and fit it accurately to the test-tube by means of a cork. (Longer tubes may be neatly cut to any size by smartly drawing the edge of a triangular file across the glass at the required point, then clasping the tube, the scratch being between the hands, and pulling the portions asunder, force being exerted in a *slightly* curved direction so as to open out the crack which the file has commenced.) The tube is fixed in the cork through a round hole made by the aid of a red-hot wire, or, better, a rat-tail file, or, best of all, by one of a set of cork-borers—pieces of brass tubing sharpened at one end and having a flat head at the other. Setting aside the test-tube for a few minutes, proceed to bend the long piece of tubing to the most convenient shape for collecting the gas.

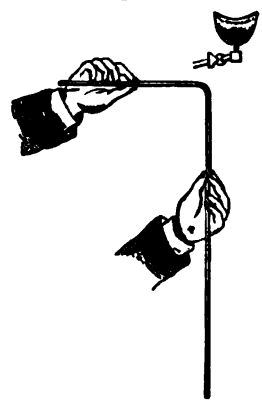
*To Bend Glass Tubes.*—Hold the part of the tube required to be bent in any gas- or spirit-flame (a fish-tail gas-jet answers

Fig. 1.

Softening and bending Glass Tubes.

very well), constantly rotating it, so that about an inch of the glass becomes heated. It will soon be felt to soften, and will then, yielding to the *gentle* pressure of the fingers, assume any required angle. In the present case, the tube should be heated at about four inches from the extremity to which the cork is attached, and bent to an angle of about 90 degrees.

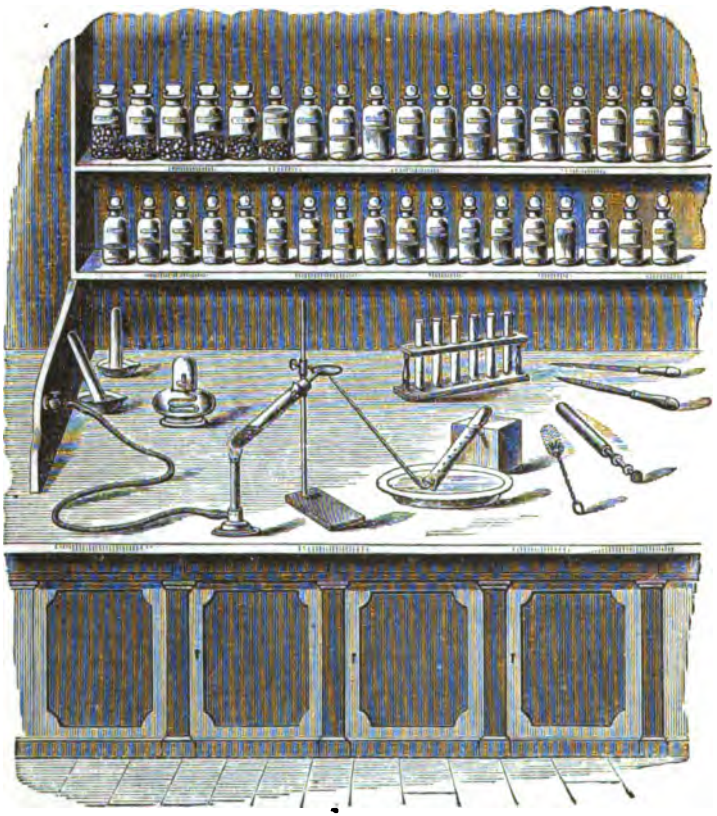
Fig. 2.



*Source of Heat.*—The source of heat for the test-tube may be the flame of an ordinary spirit-lamp, or, still better where coal-gas is procurable, a mixture of the latter with air. Gas-lamps especially constructed to burn a mixture of coal-gas and air are sold by chemical-apparatus manufacturers. (See Figs. 3 and 7).

*Collection, etc.* (*continued*).—Fit the cork and bent tube into the test-tube; the apparatus will then be ready for delivering gas at a convenient distance from the heated portion of the arrangement. To collect it, have ready three or four test-tubes (or small wide-mouthed bottles) filled with water, and inverted in a basin, or other similar vessel, also containing water, taking care to keep the mouths of the tubes a little below the surface. Now apply heat to the chlorate contained in the test-tube, and so arrange the open end of the bent tube under the water that the gas which presently issues may bubble into and gradually fill the inverted test-tubes. The first tubeful may be rejected, as it probably consists of little more than the air originally in the apparatus, and which has been displaced by the oxygen. That which comes afterwards will be pure oxygen.

Fig. 3.



This engraving represents the preparation, collection, and storage of small quantities of oxygen gas. A test-tube and bent glass tube, joined together by a perforated cork, are supported by the arm of an iron stand. (The apparatus might be held by the fingers.) The tube is heated by a gas-lamp. (The spirit-lamp shown at the back might be used instead.) Gas evolved from the heated substance in the test-tube is displacing water from an inverted test-tube. Spare tubes in a test-tube rack are at hand, and tubes already filled are set aside till wanted. A nest of cork-borers, a round file, a triangular file, and a test-tube cleaning brush are lying on the table or student's bench. Below are cupboards for apparatus, above are bottles containing testing liquids, etc.

As each tube- or bottleful is collected, the mouth (still under the surface of the water) may be closed by a cork and set aside; or a little cup (such as a porcelain crucible or small gallipot) may be brought under the mouth, and the cup, with the mouth of the tube in it, be lifted out of the water and placed close by till wanted, the water remaining in the cup effectually preventing the gas from escaping.

*On the large scale*, oxygen may be made in the same way, larger vessels (glass flasks or iron bottles) being employed. Less heat also will be necessary if the chlorate of potassium be previously mixed with very fine sand, or, still better, with about an equal weight of common black oxide of manganese.

*Note on the Collection and Storage of Gases.*—It may be as well to state that nearly all gases, whether for experimental or practical purposes, are collected and stored in a similar manner. Even coal-gas is generated at gas-works in iron retorts very much the shape of test-tubes, only they are as many feet long as a test-tube is inches; and the well-known gigantic gas-holders may be viewed as inverted iron test-tubes of great diameter.

*Properties.*—Oxygen is a colorless gas. Cailletet and Pictet have succeeded in liquefying it. Obviously it is not very soluble in water, or it could not be collected by the aid of that liquid.

Oxygen is soluble to a certain extent, however (about 3 volumes in 100 at common temperatures), or fishes could not breathe.

Other noticeable features are its want of taste and smell. Next, to show the relation of oxygen to combustion, remove one of the tubes from the water by placing the thumb over its mouth, apply for a second a lighted wood match to the orifice; the gas will be found to be incombustible. Extinguish the flame of the match, and then quickly introduce the still incandescent carbonaceous extremity of the wood halfway down the test-tube; the wood will at once burst into flame owing to the extreme violence with which oxygen supports combustion. These tests of the presence of oxygen may also be applied at the extremity of the delivery-tube whilst the gas is being evolved. (It is desirable to retain two tubes of the gas for use in subsequent experiments; also one tube in which only one-third of the water has been displaced by oxygen.)

*Relation of Oxygen to Animal and Vegetable Life.*—Not only the carbon at the end of a piece of charred wood, but any other substance that will burn in air (which, as will be seen presently, is diluted oxygen) will burn more brilliantly in pure oxygen. The warmth of the body of animals is kept up by the continuous burning of the tissues

in the oxygen (of the air) drawn into the system through the lungs. The product of this combustion is a gaseous compound of carbon and oxygen termed carbonic acid gas, a gas which, in sunlight, is decomposed in the cells of plants, with fixation of the carbon and liberation of the oxygen; hence the atmosphere is kept constant in composition.

*Memorandum.*—At present it is not advisable that the reader should trouble himself with the consideration of the chemical action which occurs either in the elimination of oxygen from its compounds, or in the separation of any of the following non-metallic elements from their combinations. It is to the properties of the elements themselves that he should restrict his attention. Working thus from simple to more complex facts, he will in due time find that the comprehension of such actions as occur in the preparation of these few elements will be easier than if he attempted their full study now.

## HYDROGEN.

*Preparation and Collection.*—The element hydrogen is also a gas,* and is obtainable from its commonest compound, water (of which one-ninth by weight is hydrogen), by the agency of hot zinc or iron, but more conveniently by the action of either of these metals on cold diluted sulphuric acid. The apparatus used for making oxygen may be employed for this experiment; but no lamp is required. Place several pieces of thin zinc† in the generating-tube (Fig. 4), or in any common glass bottle (Fig. 5) or flask, and cover them with water. The collecting tubes (these may be wide-mouthed bottles) being ready, add strong sulphuric acid (oil of vitriol) to the zinc and water, in the proportion of about one volume of acid to five of water, and fit on the delivery-tube, or pour the acid down such a funnel-tube‡ as is shown in Fig. 5; the hydrogen is at once

* Graham obtained alloys of hydrogen with palladium and other metals, compounds in which several hundred times its bulk of gas is retained by the metal in vacuo or even at a red heat. This was physical confirmation of the opinion long held by chemists, that hydrogen is a gaseous metal. Graham termed it hydrogenium (other chemists *hydrium*), and considered its relative weight in the solid state to be nearly three-fourths that of water. Cailletet and Pictet have since actually liquefied and solidified this element.

† The best form is *granulated zinc* (*Zincum Granulatum*, B P.), made by heating scraps of common sheet zinc in a ladle over a fire, and as soon as melted pouring, in a slow stream, into a pail of water from a height of 8 or 10 feet. Each drop of zinc thus yields a thin little bell, which, for its weight, presents a large surface to the action of the acid water. If the zinc is allowed to become hotter than necessary, the little bells will not be formed.

‡ Funnel-tubes may be purchased of the apparatus-maker, or, if the pupil has access to a table blowpipe and the advantage of a tutor to direct his operations, they may be made by himself.

evolved. Having rejected the first portions (or having waited until the air originally in the bottle may be considered to be all expelled), collect four or five tubes of the gas in the manner described under Oxygen.

Fig. 4.

Fig. 5.

Preparation of Hydrogen.

In making larger quantities bottles of appropriate size may be employed.

Other metals, notably potassium and sodium, liberate hydrogen the moment they come into contact with water; but the processes are not economical.

*Properties.*—Like oxygen, hydrogen is invisible, inodorous, and tasteless. If made with iron it has a strong smell, but this is due to impurities contained in the metal.

Apply a flame to the mouth of the delivery tube (care being taken that the gas is coming off briskly—a guarantee that no air remains in the generating vessel); ignition of the hydrogen ensues, showing that, unlike oxygen, it is combustible.

Immerse a lighted match into a tube (or wide-mouthed bottle) containing hydrogen; the gas is ignited, but the match becomes extinguished. This shows that hydrogen is not a supporter of combustion.

Hydrogen in burning unites with the oxygen of the air and forms water, which may be condensed on a cool glass or other surface. Prove this by holding a glass vessel a few inches above a hydrogen flame. In burning the hydrogen contained in one of the tubes or bottles, the flame is best seen when the tube is held mouth upwards, and water poured in so as to force out the gas gradually.

If, instead of this gradual combination of the two elements oxygen and hydrogen, they be mixed together in bulk in the

right proportions and then ignited, they will rapidly combine, and explosion will result. Prepare a mixture of this kind by filling up with hydrogen a test-tube from which one-third of the water has been expelled by oxygen. Remove the tube from the water, placing a finger over the mouth, and, having a lighted match ready, apply the flame; a slight explosion ensues, owing to the instantaneous combination of the two elements, and the expansive force of the highly heated steam produced. If anything larger than a test-tube is employed in this experiment, it should be a soda-water bottle, or some such vessel equally strong.

These two gases thus unite at a temperature far higher than that of boiling water, two volumes of hydrogen and one of oxygen yielding two volumes of gaseous water (true steam).

*The noise of such explosions* is caused by concussion between the suddenly expanded gaseous body and the air.

*The force of the explosion*, or, in other words, the expansive force of the suddenly heated, and therefore suddenly expanded, steam, is below that necessary to break the test-tube. Some force, however, is exerted, and hence the necessity of the precaution previously suggested of allowing all the air which may be in a hydrogen-apparatus to escape before proceeding with the experiments. If a flame be applied to the delivery-tube before all the air is expelled, the probable result will be ignition of the mixture of hydrogen and oxygen (of the air) and consequent explosion. But even in this case the generating-vessel is not often fractured unless it be large and of thin glass, the ordinary effect being that the cork is blown out, and the delivery-tube broken on falling to the ground.

Hydrogen is a prominent constituent of all the substances used for producing artificial light, such as tallow, oil, and coal-gas. The explosive force of large quantities, such as a roomful of coal-gas and air, though vastly below that of an equal weight of gunpowder, is well known to be sufficient at least to blow out that side of the room which offers least resistance.

*The composition of water* can be proved analytically as well as synthetically, a current of electricity decomposing it into its constituent gases, twice as much hydrogen as oxygen, by volume, being produced.

*Combustion* (from *comburo*, to burn).—The experiments with hydrogen and oxygen illustrate the true character of combustion. Whenever chemical combination is sufficiently intense to be accompanied by heat and light, the materials are said to undergo combustion. Combustion only occurs at the line of contact of the combining bodies; a jet of oxygen will burn in an atmosphere of hydrogen quite as easily as a jet of hydrogen in oxygen. A jet of air (diluted oxygen) will burn as readily in a jar of coal-gas as a jet of coal-gas burns in air; each is combustible, each supports the combustion of the other. Hence the terms *combustible* and *supporter of combustion* are purely conventional, and only applicable so long as the cir-

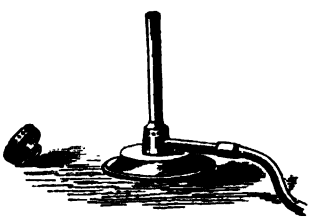
cumstances under which they are applied remain the same. In the case of substances burning in air, the conditions are, practically, always the same: hence no confusion arises from regarding air as the great supporter of combustion, and bodies which burn in it as being combustible.

*Structure of Flame.*—A candle-flame or oil-flame is a jet of gas intensely heated; the central portion is unburnt gas; the next envelope is formed of partially burnt and very dense, gaseous, and solid particles heated sufficiently high to give light, and the outer cone of completely burnt gases. In the figure the sharpness of limit of these cones is purposely somewhat exaggerated. Air made, by any mechanical contrivance of burner, to mix with the interior of a flame at once burns up, or perhaps prevents the formation of dense gases giving a hotter, but non-luminous, jet. The air-gas lamps (Fig. 7), or "Bunsen" gas-burners commonly used in chemical laboratories are constructed on this principle; their flame has the additional advantage of not yielding a deposition of soot.

Fig. 6.

Structure of Flame.

Fig. 7.



"Bunsen" or Air-gas Burner.

In the air-gas lamp, coal-gas escaping from a small orifice draws rather more than twice its volume of air (supplied through adjacent holes) into its column, and the mixture of gas and air passes upwards along a pipe. It only burns at the end, and not within the pipe, partly because the metal of the burner, by conducting heat away, cools the mixture below the temperature at which it can ignite; partly because the velocity with which the mixture flows out is greater than the rate at which such a mixture ignites; and partly because the proportion of air to gas in this mixture is insufficient for thorough and perfect combustion, the external air contributing materially to the complete combustion of the jet of air-gas. The *Davy safety-lamp* acts on the first-named principle; a wire-gauze cage surrounds an oil-flame; an inflammable mixture of gas and air (fire-damp) can pass through the gauze and catch fire and burn inside; but the flame cannot be communicated to the mixture outside, because the metal of the gauze and of the other parts cools down the gas below the temperature at which combustion can continue.

*Properties* (*continued*).—Hydrogen is the lightest substance known. It was formerly used for filling balloons, but was soon superseded by coal-gas. Coal-gas is not so light as hydrogen, but is cheaper and more easily obtained. The lightness of hydrogen may be rendered evident by the following experiment: Fill two test-tubes with the gas, and hold one with its mouth downwards and the other with its mouth upwards. The hydrogen will have escaped from the latter in a few seconds, whereas the former will still contain the gas after the lapse of some minutes. This may be proved by applying a lighted match to the mouths of the respective tubes.

*The relative weight* or specific gravity of oxygen is sixteen times that of hydrogen. A vessel holding one grain of hydrogen will hold sixteen grains of oxygen. The relation of the weight of hydrogen to air is as 1 to 14.44 or as 0.0693 to 1.0. One grain of hydrogen by weight would measure about 27 fluidounces. One grain of hydrogen would, therefore, about fill a common English wine bottle. Such a bottle would hold about 14½ grains of air, or 16 grains of oxygen.

*Mem.*—It is desirable to retain two tubes of hydrogen for use in subsequent experiments.

*Diffusion of Gases.*—Hydrogen cannot be kept in such vessels as the inverted test-tubes; for, though much lighter than air, it *diffuses* downwards into the air, while the air, though much heavier, diffuses upwards into the hydrogen. This power of *diffusion* is characteristic of all gases, and proceeds according to a fixed law, namely, "in inverse proportion to the square root of the specific gravity of the gas" (Graham). Thus hydrogen diffuses four times faster than oxygen. This great and important property of diffusion strongly suggests that the particles of gases, at least, are always moving, never at rest; how otherwise *could* gases diffuse into each other, as they do, notwithstanding the opposing influence of gravitation? Diffusion strongly supports this (Clausius's) Kinetic (*κινέω*, *kineō*, I move or put in motion) theory of the physical condition of gases.

## PHOSPHORUS.

*Appearance and Source.*—Phosphorus (*Phosphorus*, B. P. and U. S. P.) is a solid element, in appearance and consistence resembling white wax; but it gradually becomes yellow by exposure to light. It is a characteristic constituent of bones, and is always prepared from that source by a process which will be subsequently described.

*Caution.*—Phosphorus, on account of its great affinity for oxygen, takes fire very readily in the air, and should therefore be kept under water. When wanted for use it must be cut under water. It is employed in tipping lucifers, though *red* or *amorphous phosphorus* (*vide* Index) is least objectionable for this purpose.

*Properties.*—Dry a piece about one-fourth the size of a pea by quickly and carefully pressing it between the folds of porous

(filter or blotting) paper; place it on a plate, and ignite by touching it with a piece of warm wire or wood. The product of combustion is a dense white suffocating smoke, which must be confined at once by placing an inverted tumbler, test-glass, or other similar vessel over the phosphorus. The fumes rapidly aggregate, and fall in white flakes on the plate. When this has taken place, and the phosphorus is no longer burning, moisten the powder with a drop or two of water, and observe that some of the water is converted into steam, an effect due to the intense affinity with which the two combine.

The powder produced by the combustion of phosphorus is phosphoric anhydride; the combination of the latter with the elements of water produces a variety of phosphoric acid which dissolves in the water, forming on standing a dilute solution of ordinary phosphoric acid. The Diluted Phosphoric Acid of the British and United States Pharmacopœias is a somewhat similar solution, made, however, in a different way, and of a definite strength.

## NITROGEN.

*Source.*—The chief source of this gaseous element is the atmosphere, nearly four-fifths of which consists of nitrogen (the remaining fifth being almost entirely oxygen.)

*Preparation.*—Burn a piece of dried phosphorus, the size of a pea, in a confined portion of air. The oxygen is thus removed, and nitrogen alone remains. The readiest mode of performing this experiment is to fix a piece of earthenware

Fig. 8.

Preparation of Nitrogen.

Fig. 9.

Decantation of Gases.

(the lid of a small porcelain crucible answers very well) on a thin piece of cork, so that it may float in a dish of water. Place the phosphorus on the lid, ignite by a warm rod, and then in-

vert a tumbler, or any glass vessel of about a half-pint capacity, over the burning phosphorus, so that the glass may dip into the water. Let the arrangement rest for a short time for the fumes of phosphoric anhydride to subside and dissolve in the water, and then decant the gas into test-tubes in the manner indicated in Fig. 9, using a tub or other vessel of water of sufficient depth to permit the glass containing the nitrogen to be turned on one side without air gaining access.

Larger quantities of nitrogen are made in the same way. Other combustibles, such as sulphur or a candle, might be used to burn out the oxygen from a given quantity of air, but none answer so quickly and completely as phosphorus; added to which, the product of their combustion would not always be dissolved by water, but would remain with and contaminate the nitrogen.

*Mem.*—The statement concerning the composition of the air is roughly confirmed in preparing nitrogen, about one-fifth of the volume of the air originally in the glass vessel having disappeared, its place being occupied by water from the dish.

*Properties.*—Like oxygen and hydrogen, nitrogen gas is invisible, tasteless, and inodorous. By pressure Cailletet and Pictet have succeeded in condensing it to a liquid. It is only slightly soluble in water. Free nitrogen is distinguished from all other gases by the absence of any characteristic or positive properties. Apply a flame to some contained in a tube; it will be found to be incombustible. Immerse a lighted match in the gas; the flame is extinguished, showing that nitrogen is a non-supporter of combustion.

The chief office of nitrogen in the air is to dilute the energetic oxygen, a mere *mechanical* mixture resulting.

Nitrogen is fourteen times as heavy as hydrogen.

The air is nearly fourteen and a half (14.44) times as heavy as hydrogen. Its average composition, including minor constituents, which will be referred to subsequently, is as follows:—

*Composition of the Atmosphere.*

| | In 100 volumes. |
|---|---|
| Oxygen | 20.61 |
| Nitrogen | 77.95 |
| Carbonic acid gas | .04 |
| Aqueous vapor | 1.40 |
| Nitric acid, Ammonia, Carburetted hydrogen | traces. |
| Sulphuretted hydrogen, Sulphurous acid | traces in towns. |

The above proportions are by volume. By weight there will be nearly 23 parts of oxygen to nearly 77 of nitrogen, oxygen being

the heavier in the ratio of 16 to 14. Ozone (*vide* Index) is also said to be a normal constituent of air.

The comparative inactivity or negative character of nitrogen in its free condition, that is, when uncombined with other elements, contrasts strongly with its apparent influence in a state of combination. When its compounds with hydrogen come to be studied, it will be found to be, apparently, the chief, or leading, or, in a sense, the most important element of those compounds—the ammoniacal compounds. United with carbon it gives the poisonous cyanic substances. With oxygen it gives quite a large group of bodies, amongst which are the common and important class of salts termed nitrates. With carbon as well as hydrogen and some oxygen it affords powerful agents termed alkaloids—near relatives of ammonia—while the same elements otherwise grouped, with sometimes a little sulphur or phosphorus, form the various albumenoid and gelatinous matters characteristic of the tissues of animals and vegetables. In a perfect structure we should perhaps scarcely regard any one element or member as more important than another, still such a conclusion almost forces itself upon us as we become acquainted with the chemical history of combined nitrogen.

## CHLORINE.

*Source.*—This element is a gas. Its chief source is common salt, more than half of which is chlorine.

*Preparation.*—About a quarter of an ounce of salt and the same amount of black oxide of manganese are mixed, and placed in a test-tube with sufficient water to cover them; on adding a small quantity of sulphuric acid, the evolution of chlorine commences. For mode of collection see following paragraphs.

Fig. 10.

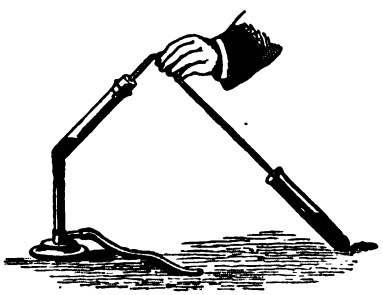


Fig. 11.

Preparation of Chlorine.

*Another Process.*—As the action of the sulphuric acid on the salt in the above process is mainly to give hydrochloric acid, the latter

acid (about 4 parts) and the black oxide of manganese (about 1 part) may be used in making the gas, instead of salt, sulphuric acid, and black oxide of manganese. This, the usual process, is that adopted in the British and United States Pharmacopœias.

*Collection and properties.*—Chlorine is a most suffocating gas. Great care must consequently be observed in experimenting with this element. As soon as its penetrating odor indicates that it is escaping from the test-tube, the cork and delivery-tube (similar to that used in making oxygen) should be fitted on, and the gas allowed to pass to the bottom of another test-tube containing water (Fig. 10). When thirty or forty small bubbles have passed, their evolution being assisted by slightly heating the generating-tube, the latter should be removed to the cupboard usually provided in laboratories for performing operations with noxious gases, or dismounted, and the contents washed away. The water in the collecting-tube will now be found to smell of the gas, chlorine being, in fact, soluble in about half its bulk of water. Chlorine-water is official* in the British and United States Pharmacopœias (*Liquor Chlori*, B. P., *Aqua Chlorinii*, U. S. P.).

*Larger quantities* may be made from hydrochloric acid and black oxide of manganese (about 4 parts to 1) in a Florence flask, fitted with a delivery-tube, the flask being supported over a flame by the ring of a retort stand or any similar mechanical contrivance (Fig. 11). A piece of cardboard on the neck of the collecting-bottle, as indicated in the figure, retards diffusion of the gas from the bottle during collection of the gas.

*Mem.*—Flasks and similar glass vessels are less liable to fracture if protected from the direct action of the flame by being placed on a piece of wire gauze 3 to 4 inches square, or on a *sand-bath*, that is, a saucer-shaped tray of sheet iron, on which a *thin* layer of sand is placed.

---

* The Pharmacopœia and all in it are official (*office*, Fr., from L. *officium*, an office). There are many things which in pharmacy are officinal (Fr., from L. *officina*, a shop) but not official. To restrict the word *officinal* to the contents of a pharmacist's shop, and to that portion of the contents which is Pharmacopœial, is radically wrong, and should be avoided. "An *official* formula is one given under authority. An *officinal* formula is one made in obedience to the customary usage of the shop (*officina*). To state that any preparation under the sanction of the Pharmacopœia is officinal, is a misapprehension of the meaning of the word."—*J. Brough.*

That is *official* which emanates from a recognized authority. That is *officinal* which is issued from an *officina* or workshop.—*Joseph Ince.*

*Official* writings and orders are those issued by official persons. *Officinal* articles are such as are found in a shop.—*J. F. Stanford, M.A., F.R.S.*

The *Vapor Chlori*, B. P., or Inhalation of Chlorine, is simply moist chlorinated lime so placed that some of the chlorine given off may be inhaled.

During these manipulations the operator will have noticed that chlorine is of a light green color. That tint is readily observed when the gas is collected in large vessels. As it is soluble in water 2½ vols. in 1 vol. at 60° F.), it cannot be economically stored over that liquid. Being, however, nearly twice and a half as heavy as air, it may be collected by simply allowing the delivery tube to pass to the bottom of the test-tube or dry bottle (Fig. 11).

The distinctive property of chlorine is its bleaching-power. Prepare some colored liquid by placing a few chips of logwood or other dyeing material in a test-tube half full of hot water. Pour off some of this red decoction into another tube and add a few drops of the chlorine-water; the red color is rapidly destroyed.

Chlorine readily decomposes offensive effluvia; it is one of the most powerful of the *deodorizers*. It also decomposes putrid and infectious matter; it is one of the best of *disinfectants*. (*Antiseptics* are substances which *prevent* putrefaction. See Index.)

*Combination of Hydrogen with Chlorine, forming Hydrochloric Acid.*—If an opportunity occurs of generating the gas in a closed chamber or in the open air, a test-tube, of the same size as one of those in which hydrogen has been retained from a previous operation, is filled with the gas. The hydrogen tube is then inverted over that containing the chlorine, the mouths being kept together by encircling them with a finger. After the gases have mixed, the mouths of the tubes are quickly in succession brought near a flame, when explosion occurs, and fumes of a compound of hydrochloric acid with the moisture of the air are formed. The Hydrochloric Acid of Pharmacy (*Acidum Hydrochloricum*, B. P., *Acidi Muriaticum*, U. S. P.) is a solution of the gas (made in a more economical way) in water.

The foregoing experiment affords evidence of the powerful affinity of chlorine and hydrogen for each other. Chlorine dissolved in water will, in sunlight, slowly remove hydrogen from some of the water and liberate oxygen. The bleaching power of chlorine is generally referred to this oxidizing effect which it produces in presence of water; for dry chlorine does not bleach.

*Density.*—Chlorine is thirty-five and a half times as heavy as hydrogen. A wine bottle would hold about 35½ grains.

## SULPHUR, CARBON, IODINE.

*The physical properties* of these elements (color, hardness, weight, etc.) are familiar. Their leading chemical characters will also be understood when a few facts concerning each are made the subject of experiment.

SULPHUR.—Burn a small piece of sulphur; a penetrating odor is produced, due to the formation of a colorless gas, the same as that formed on igniting a sulphur-tipped lucifer match.

This product is a perfectly definite chemical compound of the oxygen of the air with the sulphur. It is termed sulphurous anhydride or sulphurous acid gas.

CARBON is familiar in the forms of soot, coke, charcoal, graphite (or plumbago, popularly termed blacklead), and diamond. The presence of carbon in wood, and in other vegetable and animal matter, is at once rendered evident by heat. Place a little tartaric acid on the end of a knife in a flame; the blackening that occurs is due to the separation of carbon. The black matter at the extremity of a piece of half-burned wood is also carbon.

Carbon, like hydrogen, phosphorus, and sulphur, has a great affinity for oxygen at high temperatures. A striking evidence of that affinity is the evolution of sufficient heat to make the materials concerned red or even white hot. When ignited in the dilute oxygen of the air, carbon simply burns with a moderate glow, as seen in an ordinary coke or charcoal fire, but when ignited in pure oxygen, the intensity of its combination is greatly exalted. The product of the combination of the two elements, if the oxygen be in excess, is an invisible gaseous body termed carbonic acid gas; if the carbon be in excess, another invisible gas termed carbonic oxide results.

IODINE.—A prominent chemical characteristic of iodine is its great affinity for metals. Place a piece of iodine, about the size of a pea, in a test tube with a small quantity of water, and add a few iron-filings or small nails. On gently warming this *mechanical* mixture, or even shaking if longer time be allowed, the color and odor of the iodine disappear; it has *chemically* combined with the iron; a *chemical compound* has been produced. If the solution be filtered, a clear aqueous solution of the compound of the two elements is obtained.

This compound is an iodide of iron. Its solution, made as above, and mixed with sugar, forms, when of a certain strength, the ordinary Syrup of Iodide of Iron of pharmacy (*Syrupus Ferri Iodidi*, B. P. and U. S. P.). A strong solution mixed with sugar and liquorice-root (sugar, liquorice, liquorice-root, gum Arabic, and reduced iron,

U. S. P.) constitutes the corresponding Pill (*Pilula Ferri Iodidi*, B. P. and U. S. P.). The solid iodide (*Ferri Iodidum*, B. P.) is obtained on removing the water of the above solution by evaporation.

*Sulphur* and *Iron*, also, when very strongly heated, *chemically combine* to form a substance which has none of the properties of a *mixture* of sulphur and iron, that is, has none of the characters of sulphur and none of iron, but new properties altogether. The product is termed Sulphide of Iron. Its manufacture and uses will be alluded to in treating of the compounds of iron; it is mentioned here as a simple but striking illustration of the difference between a *chemical compound* and a *mechanical mixture*.

## THE ELEMENTS, THEIR SYMBOLS, ETC.

From the foregoing statements a general idea will have been obtained of the nature of several of the more frequently occurring elements. Some additional facts concerning them may be gathered from the following Table, which gives the name in full, the symbol (or short-hand character*) of the name, and its origin.

For the purposes of study the elements may be divided into three classes, viz., those frequently used in pharmacy, those seldom, and those never used.

| Name. | Symbol. | Derivation of Name. |
|---|---|---|
| Oxygen | O | From ὀξύς (oxŭs) *acid*, and γίνεσις (genesis) *generation*, i. e., generator of acids. It was supposed to enter into the composition of all acids when first discovered. |
| Hydrogen | H | From ὕδωρ (hudōr) *water*, and γίνεις, (genesis) *generation*, in allusion to the product of its combination in air. |
| Nitrogen | N | From νίτρον (nitron), and γίνεσις (genesis), *generator of nitre.* |
| Carbon | C | From *carbo*, *coal*, which is chiefly carbon. |
| Chlorine | Cl | From χλωρὸς (chlōros) *green*, the color of this element. |
| Iodine | I | From ἰὸν (ion) *a violet*, and εἶδος (eidos) *likeness*, in reference to the color of its vapor. |
| Sulphur | S | From *sal* a *salt*, and πῦρ (pūr) *fire*, indicating its combustible qualities. Its common name, *brimstone*, has the same meaning, being the slightly altered Saxon word *brynstone*, i. e., burnstone. |
| Phosphorus | P | φῶς (phōs) *light*, and φέρειν (pherein) *to bear*. The light it emits may be seen on exposing it in a dark room. |

* The symbol is also much more than the short-hand character, as will be presently apparent.

| Name. | Symbol. | Derivation of Name. |
|---|---|---|
| Potassium ..... (Kalium.) | K | *Kalium*, from *kali*, Arabic for *ashes*. Manufactories in which certain compounds of potassium and allied sodium salts are made are called alkali-works to this day. *Potassium*, from *pot-ash;* so called because obtained by evaporating the lixivium of wood-ashes in pots. From such ashes the element was first obtained, hence the name. |
| Sodium..... (Natrium.) | Na | *Natrium*, from *natron*, the old name for certain natural deposits of carbonate of sodium. *Sodium*, from *soda-ash* or *sod-ash*, the residue of the combustion of masses or *sods* of marine plants. These were the sources of the metal. |
| Ammonium..... | Am ($NH_4$) | This body is not an element; but its components exist in all ammoniacal salts, and apparently play the part of such elements as potassium and sodium. Sal ammoniac (chloride of ammonium) was first obtained from near the temple of Jupiter Ammon in Libya; hence the name. |
| Barium..... | Ba | From βαρὺς (barûs) *heavy*, in allusion to the high specific gravity of "heavy spar," the most common of the barium minerals. |
| Calcium..... | Ca | *Calx*, *lime*, the oxide of calcium. |
| Magnesium ..... | Mg | From *Magnesia*, the name of the town (in Asia Minor) near which the substance now called "native carbonate of magnesia" was first discovered. |
| Iron ..... (Ferrum.) | Fe | The spelling is from the Saxon *iren*, the pronunciation probably from the kindred Gothic "*iarn;*" the derivation is Aryan; it probably originally meant *metal*. |
| Aluminium ..... | Al | The metallic basis of alum was at first confounded with that of sulphate of iron, which was the alum of the Romans, and was so called in allusion to its tonic properties, from *alo*, to *nourish*. |
| Zinc..... | Zn | From Ger. *zinn*, *tin*, with which zinc seems at first to have been confounded. |
| Arsenicum ..... | As | Ἀρσενικον (arsenikon), the Greek name for orpiment, a sulphide of arsenicum Common white arsenic is an oxide of arsenicum. |
| Antimony..... (Stibium.) | Sb | στίβι (stibi), or στίμμι (stimmi), was the Greek name for the native sulphide of antimony. The word *antimony* is said to be derived from ἀντὶ (anti *against*), and *moine*, French for *monk*, from the fact that certain monks were poisoned by it. |
| Copper..... (Cuprum.) | Cu | From *Cyprus*, the name of the Mediterranean island where this metal was first worked. |

| Name. | Symbol. | Derivation of Name. |
|---|---|---|
| Lead..... (Plumbum.) | Pb | The Latin word is expressive of "something heavy," and the Saxon *læd* has a similar signification. |
| Mercury ..... (Hydrargyrum.) | Hg | *Hydrargyrum*, from ὕδωρ (hudōr) *water*, and ἄργυρος (arguros) *silver*, in allusion to its liquid and lustrous characters. *Mercury*, after the messenger of the gods, on account of its susceptibility of motion. The old name *quicksilver* also indicates its ready mobility and argentine appearance. |
| Silver..... (Argentum.) | Ag | Ἄργυρος (arguros) *silver*, from ἀργὸς (argos) *white*. Words resembling the term *silver* occur in several languages, and indicate a white appearance. |

The following are the names of some of the less frequently occurring elements, compounds of which, however, are alluded to in the British and U. S. Pharmacopœias, or met with in pharmacy.

| Name. | Symbol. | Derivation of Name. |
|---|---|---|
| Bromine ..... | Br | From βρῶμος (brōmos), a *stink*. It has an intolerable odor. |
| Fluorine | Fl | Fluo, *to flow*. Fluoride of calcium, its source is commonly used as a flux in metallurgic operations. |
| Boron..... | Bo | From *borax*, or *baurak*, the Arabic name of *borax*, the substance from which the element was first obtained. |
| Silicon ..... | Si | From *silex*, Latin for *flint*, which is nearly all silica (an oxide of silicon). |
| Lithium..... | L | From λίθιος (litheios) *stony*, in allusion to its supposed existence in the mineral kingdom only. |
| Strontium ..... | Sr | This name is commemorative of *Strontian*, a mining village in Argyleshire, Scotland, in the neighborhood of which the mineral known as *strontianite* or carbonate of strontium was first found. |
| Cerium..... | Ce | Discovered in 1803, and named after the planet *Ceres*, which was discovered on Jan. 1, 1801. The oxalate of cerium is official, but seldom used. |
| Chromium..... | Cr | From χρῶμα (chrōma) *color*, in allusion to the characteristic appearance of its salts. |

| Name. | Symbol. | Derivation of Name. |
|---|---|---|
| Manganese ..... | Mn | Probably a mere transposition and repetition of most of the letters of the word magnesia, with whose compounds those of manganese were confounded till the year 1740. |
| Cobalt ..... | Co | *Cobalus* or *Kobold* was the name of a demon supposed to inhabit the mines of Germany. The ores of cobalt were formerly troublesome to the German miners, and hence received the name their metallic radical now bears. |
| Nickel..... | Ni | *Nickel*, from *nil*, is a popular German term for *worthless*. The mineral now known as nickel ore was formerly called by the Germans *Kupfernickel*, *false copper*, on account of its resemblance to copper (*Kupfer*) ore. When a new metallic element was found in the ore, the name nickel was retained. |
| Tin (Stannum)... | Sn | Both words are possibly corruptions of the old British word *staen*, or the Saxon word *stan* a stone. Tin was first discovered in Cornwall, and the ore (an oxide) is called *tinstone* to the present day. |
| Gold (Aurum) .... | Au | *Aurum* (Latin) from a Hebrew word signifying the color of fire.<br>*Gold*. A similar word is expressive of *bright yellow* in several old languages. |
| Platinum ..... | Pt | From *platina* (Spanish) diminutive of *plata*, *silver*. It somewhat resembles silver in appearance, but is less white and lustrous. |
| Bismuth ..... | Bi | Slightly altered from the German *Wismuth*, derived from *Wiesematte* "a beautiful meadow," a name given to it originally by the old miners in allusion to the prettily variegated tints presented by the freshly exposed surface of this crystalline metal. |
| Cadmium ..... | Cd | Καδμεία (Kadmeia) was the ancient name of calamine (carbonate of zinc), with which carbonate of cadmium was long confounded, the two often occurring together. |

Gold, Platinum, Tin, and Silicon are classed with the less important elements, because their salts are seldom used in pharmacy.

It will be noticed that the symbol of an element is simply the first letter of its Latin name, which is generally the same as in the English. Where two names begin with the same letter, the less important has an additional letter added.

## QUESTIONS AND EXERCISES.

1. Of how many elements is terrestrial matter composed?
2. In what state do the elements occur in nature?
3. Distinguish between the *art* and the *science* of chemistry.
4. What is the difference between an element and a compound?
5. Enumerate the chief non-metallic elements.
6. Describe a process for the preparation of oxygen.
7. How are gases usually stored?
8. Mention the chief properties of oxygen.
9. What is the source of animal warmth?
10. State the proportion of oxygen in air.
11. Is the proportion constant, and why?
12. Give a method for the elimination of hydrogen from water.
13. State the properties of hydrogen.
14. Why is a mixture of hydrogen and air explosive?
15. Explain the effects producible by the ignition of large quantities of coal-gas and air.
16. What is the nature of combustion?
17. Define a *combustible* and a *supporter of combustion.*
18. Describe the structure of flame.
19. State the principle of the Davy safety-lamp.
20. To what extent is hydrogen lighter than oxygen?
21. What do you mean by *diffusion* of gases?
22. State Graham's law concerning diffusion.
23. Name the source of phosphorus, and give its characters.
24. Why does phosphorus burn in air?
25. What remains when ignited phosphorus has removed all the oxygen from a confined portion of air?
26. Mention the properties of nitrogen.
27. What office is fulfilled by the nitrogen of air?
28. State the proportions of the chief constituents of air.
29. Mention the minor or occasional constituents of air.
30. What is the proportion by weight of nitrogen to oxygen in the atmosphere?
31. Give the specific gravity of nitrogen?
32. How is chlorine prepared?
33. Enumerate the properties of chlorine.
34. Define the terms *deodorizer* and *disinfectant.*
35. Explain the bleaching effect of chlorine.
36. What proportion of hydrogen to chlorine is necessary for the formation of hydrochloric acid gas?
37. State the prominent physical and chemical characters of sulphur.
38. State the prominent characters of carbon.
39. State the prominent characters of iodine.
40. Give the derivations of the names of some of the elements.
41. What are the symbols of oxygen, hydrogen, nitrogen, carbon, chlorine, iodine, sulphur, phosphorus?

THE LEARNER IS RECOMMENDED TO READ THE FOLLOWING PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY CAREFULLY ONCE OR TWICE, THEN TO STUDY (EXPERIMENTALLY, IF POSSIBLE) THE SUCCEEDING PAGES, RETURNING TO AND READING OVER THE GENERAL PRINCIPLES FROM TIME TO TIME UNTIL THEY ARE THOROUGHLY COMPREHENDED.

## THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY.

### DEFINITION OF CHEMICAL ACTION.

The learner may now proceed to study the manner in which substances act chemically on each other. By acting *chemically* it will be obvious, from the preceding experiments, that what is meant is *so affecting each other that the substances are greatly altered in properties*. A *mixture* of oxygen and hydrogen is still a gas; *a chemical compound* of oxygen and hydrogen is a liquid, namely, water; here is great alteration in leading properties. Iodine is only slightly soluble in water, and forms a brown-colored solution, and iron is insoluble; but when iodine and iron are *chemically* combined, the product is very soluble in water, forming a light-green solution in which the eye can detect neither iodine nor iron, and which is utterly unlike iron or iodine in any one of their properties. Sand, sugar, and butter rubbed together form *a mere mixture*, from which water would extract the sugar, and ether dissolve out the butter, leaving the sand. Tartaric acid, carbonate of sodium, and water added to each other, form a *chemical compound*, containing neither tartaric acid nor carbonate of sodium, these bodies having attacked each other and formed fresh combinations. These illustrations show that *chemical* action is distinguished from all other actions by (*a*) producing an entire change of properties in the bodies on which it is exerted. Chemical action is further distinguished by (*b*) the fact that it only takes place between definite weights and volumes of matter. This (*a* and *b*) cannot be said of any other action—the action of any of the other great forces of nature (gravitation, heat, light, electricity, etc.); hence the statements (*a* and *b*) furnish a sharp and precise definition of *chemical* action or the chemical force, the force whose manifestations the reader of this book, or of any other manual of chemistry, is studying.

### ATOMS.

In a chemical compound, what has become of the constituents? Let the reader place before him specimens of sulphur, iron, and sulphide of iron; or iodine, iron, solid iodide of iron and its solution in water or syrup (*Syrupus Ferri Iodidi*, B. P. and U. S. P.). In the sulphide of iron what has become of the sulphur and of the iron from which it was made? The *mixture* of sulphur and iron in *combining* to form sulphide of iron has not lost weight, and, indeed, by certain

processes it is possible to recover its sulphur as sulphur, and its iron as iron; so that we are compelled to believe, we cannot avoid the conclusion, that sulphide of iron contains particles of sulphur and of iron. But how small must be those particles! Rub a minute fragment to dust in a mortar and place a trace of the powder under the highest power of the best microscope; no yellow particle is visible, not the minutest portion of lustrous metal, but dull-brown miniature fragments of the original mass. The elementary particles of sulphur and iron, or of the elements in any other compound (the chlorine and sodium in common salt or the iodine and iron in solution of iodide of iron), are, in short, too small to be seen. Can they be imagined? Again, no. The mind cannot conceive of a particle of anything (sulphur, iron, sulphide of iron, or what not) so small but what the next instant the imagination has divided it. Yet learner and teacher must have some common platform on which to reason and converse. The difficulty is met by speaking of these inconceivably small particles as *atoms* (ἄτομος *atomos*, invisible; from the privative α and τεμνω, *temnō*, to cut—that which is not cut or divided), an expedient suggested by our countryman Dalton at the commencement of the present century. It is an expedient not perhaps altogether satisfactory, but is the only one possible to the majority of minds in the present state of knowledge and education. We cannot speak of iodine and iron uniting lump to lump, as two bricks are cemented together or blocks of wood glued together, for such is not the kind of action. We cannot select a minute fragment of each to regard as the combining portions, for the minutest fragment we could obtain is visible, and iodide of iron contains neither visible iodine nor visible iron. And yet iodide of iron contains both iodine and iron, or, at least, a given weight of the compound is obtained from the same weight of the constituents, and the same weight of constituents is obtainable from an equal weight of the compound. We might say that molecules are concerned in the operation, but molecules means little masses of—of what? there is positively no word left with which to carry on conversation and description but *atoms*. Any other mode of treating the matter is too subjective for general employment. Moreover, any difficulty in forming a definite conception of an atom is met by regarding an atom, not necessarily as something which *cannot* be divided, but as "a particle of matter which undergoes no further division *in chemical metamorphoses*" (Kekulé). Even physicists regard atoms from much the same point of view, indeed they often speak of still larger portions of matter (molecules) as atoms, meaning thereby "something which is not divided *in certain cases that we are considering*" (Clifford).

### The Chemical Force.

What power binds the atoms of a chemical compound together in such marvellous closeness of union that in the couple or group they lose all individuality? Clearly an attractive force of enormous power, a force remotely resembling, perhaps, that which attracts a piece of iron to a magnet. Only by such an assumption can we conceive that common salt contains chlorine and a metal (sodium), or

that wood contains carbon, hydrogen, and oxygen. Were not this force thus all-powerful, the carbon in wood would show its blackness and other qualities, and the hydrogen and oxygen give indications of their gaseous and other characters. This attractive force is commonly termed the *chemical force*, sometimes *chemical affinity*. The word *chemism* has also been proposed for it, just as the magnetic force is termed magnetism, but has not been generally adopted.

### Molecules.

A single, free, uncombined atom cannot exist in a state of isolation, at common temperatures, for any appreciable length of time. For we must regard an atom as the home of an attractive force of great intensity, and the moment it is liberated from a state of combination (hydrogen from water, for instance, or chlorine from salt) —finds itself in proximity to another atom having similar desires for union, so to speak—the result is an impetuous rushing together and formation of either couples, trios, or groups, according to the nature of the atoms. It would be as difficult to conceive of separate atoms as to imagine that a strong magnet and a piece of steel could be suspended close to each other without being drawn together. It is, doubtless, possible to keep some pairs of atoms apart by the aid of heat, just as the magnet and steel may be parted by a superior amount of force, but such a condition of things is probably abnormal. These pairs and other groups of atoms are conveniently designated by the one word *molecule*, the diminutive of *mole*, a mass; literally little masses. Dissimilar kinds of atoms seem to have greater attraction for each other than similar kinds; for, first, the masses of matter met with in nature in the great majority of cases contain two or more dissimilar elements; and, secondly, at the moment certain elements are liberated from their combination, they are very specially active in combining with other, different, elements; that is to say, the chances are not equal that the liberated elements will either retain their elementary condition or combine to form compounds, but the cases in which compounds are formed are actually in great majority.

### Recapitulation.

It is desirable that the learner should here make some experiment which will serve to bring again under notice in an applied or concrete form what has just been stated respecting the substances termed *chemical compounds*, and concerning the character of that *chemical force* which resides in the *atoms* of *molecules*. The following will usefully serve this purpose; it is the process for detecting a trace of sulphurous acid in common liquid hydrochloric acid.

As already proved, hydrogen gas and chlorine gas, when united, form hydrochloric acid gas: the latter dissolved in water is the ordinary liquid of the shops termed *Hydrochloric Acid*, the *Acidum Hydrochloricum* or *Muriaticum* of Pharmacopœias. Commercial samples of this liquid not unfrequently contain as an impurity a trace of sulphurous acid gas, a body

also already mentioned and experimentally prepared—a trace too small to be detected by its odor. Obtain a specimen of common liquid hydrochloric acid containing as an impurity a trace of sulphurous acid, or adopt the more simple course of purposely adding a few drops of aqueous solution of sulphurous acid (*Acidum Sulphurosum*,* B. P. and U. S. P.) to some hydrochloric acid. (If no sulphurous acid is at hand, the object may be accomplished by pouring a quarter or half an ounce of liquid hydrochloric acid into a wide-mouthed bottle, then burning a fragment of sulphur on a wire or strip of wood inside the bottle for a few seconds, and shaking the gas and liquid together.) Pour some of the impure liquid hydrochloric acid into a test-tube, add about an equal bulk of water, and then drop in some fragments of the metal zinc. Effervescence will occur, due to the escape of inodorous hydrogen gas, together with a small quantity of a badly-smelling gas termed sulphuretted hydrogen gas. Bring the mouth of the tube under the nose; the presence of sulphuretted hydrogen will at once be recognized.

The hydrochloric acid has now been *tested* for sulphurous acid. If the experiment be performed on any commercial specimen of the acid, and a smell of sulphuretted hydrogen be observed, the operator would at once be able to state that the specimen contains sulphurous acid as an impurity.

The true explanation—using Dalton's theory of the atomic constitution of matter—the explanation of what occurs in the successive steps of the foregoing experiment is as follows:—

Hydrochloric acid is a *chemical compound* of hydrogen and chlorine. That it is a chemical compound, and not a mere mechanical mixture of hydrogen and chlorine, is shown by the fact that its properties are altogether different from the properties of its constituents. The attractive power or *chemical force* resident in the *atoms* of chlorine and of hydrogen has caused them to combine in the closest manner imaginable and form pairs of atoms or *molecules* of the chemical compound—hydrochloric acid. Zinc being introduced into the acid, and the atoms of zinc and chlorine having even still greater attraction for each other than the hydrogen for the chlorine, the zinc and chlorine atoms combine and form a new molecule (termed chloride of zinc) which remains in the liquid, while the hydrogen atoms, having the atoms of no other element to combine with, if the acid is pure, unite to form pairs, or molecules of hydrogen, and in that state escape from the vessel. If the acid be impure from the presence of sulphurous acid (sulphurous acid gas, it will be remembered, is a compound of sulphur and oxygen), some of the hydrogen atoms, at the moment of their birth, their *nascent* state (from *nascor*, to be born)—the specially active state—finding the atoms of other elements

* These aqueous solutions of acids are generally, for the sake of brevity, simply termed acids.

present, namely, the atoms of sulphur and oxygen of the sulphurous acid, combine by preference with these atoms and form new molecules, the sulphur and hydrogen forming sulphuretted hydrogen, and the oxygen and hydrogen producing water: the former escapes with the great bulk of the hydrogen, while the water remains with the water already in the vessels.

*Note.*—Ordinary hydrogen gas, that is, hydrogen not in the nascent condition, will not thus attack sulphurous acid. Doubtless the amount, or extent, of attraction of two atoms of hydrogen for one atom of, say, the sulphur in the sulphurous acid molecule is a constant amount, but the uncombined nascent atoms can, it is only fair to suppose, get much nearer to the attacked molecule than they can after they have themselves combined to form a molecule, molecules (but not atoms) having much space between them, as will be further shown almost immediately. In other words, it is probably distance which prevents an attack which would be inevitable at close quarters. These remarks apply to all similar reactions of other elements.

### Conditions and nature of the manifestation of the Chemical Force.

The exertion of chemical affinity is only possible when substances are in close contact. Thus it was necessary to bring the oxygen, hydrogen, phosphorus, chlorine, sulphur, carbon, iodine, and iron into intimate contact, in the respective experiments with those elements, before the various reactions occurred. The exact nature of these actions, as indeed of all in which substances act chemically, would seem to be an interchange, most generally a mutual one, of the atoms of which the molecules consist—a change of partners, so to speak. Thus, in the experiment in which hydrogen and chlorine gases united to form hydrochloric acid gas, a pair of atoms in a hydrogen molecule, and a pair of atoms in a chlorine molecule, finding themselves opposite to each other, change places, the atoms of each of the old molecules unlinking, so to say, and pairing off in fresh couples—as two brothers who for many years have been close companions, and two sisters similarly united, thrown freshly into each other's society soon accept new and still more congenial couplement.

$$\left\{\begin{matrix}\text{Hydrogen}\\ \text{Hydrogen}\end{matrix}\right\} \text{and} \left\{\begin{matrix}\text{Chlorine}\\ \text{Chlorine}\end{matrix}\right\} \text{become} \left\{\begin{matrix}\text{Hydrogen}\\ \text{Chlorine}\end{matrix}\right\} \text{and} \left\{\begin{matrix}\text{Hydrogen}\\ \text{Chlorine}\end{matrix}\right\}$$

Or, using the symbols of these elements instead of the full names,

H H and Cl Cl become H Cl and H Cl.

Still further economizing space and trouble, the same statement may be made in the following form:—

$H_2$ and $Cl_2$ become 2HCl.

Once more, by using the *plus* (+) instead of the words "and" or "added to," and the sign or symbol = or *equal* instead of the words "become" or "are equal to," we reach the shortest method of expressing this chemical action:—

$$H_2 + Cl_2 = 2HCl.$$

This is the form in which such an action may be expressed in the student's note-book. It is the shortest and most convenient form, and is instructive and suggestive to the mind.

## Chemical Notation.

We have thus gradually arrived at a spot in the path of chemical philosophy at which we must halt to more fully discuss the usual method of recording chemical travels. We have arrived at the subject of *chemical notation* (from *noto*, I mark), the art or practice of recording chemical facts by short marks, letters, numbers, or other signs. Already the first capital letter, or the first and one of the following small letters of the Latin names of the elements have been employed as contractions, or short-hand expressions, or *symbols* of the whole name. Thus H has been used for the word "hydrogen," and Cl for "chlorine." A second function of such a symbol is that of indicating one atom. Thus H stands not only for the word or substance "hydrogen," but for one atom of hydrogen. Large and small figures (2 or $_2$) indicate a corresponding number of atoms, the small figure only multiplying the one particular symbol to which it is attached, while a larger figure multiplies all the symbols it precedes. Thus $H_2$ means two atoms of hydrogen, and $Cl_2$ two atoms of chlorine; while 2HCl means two atoms of hydrogen and two atoms of chlorine, or, in one word, two molecules of hydrochloric acid gas. A third function of such a symbol as H or Cl is that of indicating one volume of the element in the gaseous state. Thus H, Cl, and O stand, first, for the substances named hydrogen, chlorine, and oxygen; secondly, for single atoms of hydrogen, chlorine, and oxygen. Thirdly, they represent single and equal volumes of chlorine, hydrogen, and oxygen. It will be remembered that one test-tubeful of hydrogen and an equal sized test-tubeful of chlorine were employed in a previous experiment in forming hydrochloric acid gas, HCl.

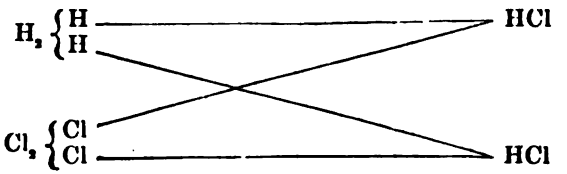
The position of symbols counts for something. Thus HCl indicates not only the substances hydrogen and chlorine, single atoms of each of the substances, and equal volumes of each, but also that the two substances are joined together by the chemical force. If the two letters were placed one under the other, or at some distance apart, or were separated by a comma or a plus sign (+), they would be understood to mean a mere mixture of the elements; but placed as close as the printer's types will conveniently and consistently allow, they must be considered to stand for a compound of the elements, that is to say, hydrochloric acid gas (HCl). The collection of symbols representing a molecule is termed a *formula*. $H_2$, $Cl_2$, and HCl are the *formulæ* of hydrogen, chlorine, and hydrochloric acid gas.

$$H_2 + Cl_2 = 2HCl.$$

Such a set of letters, figures, and marks as that on the above line is collectively termed an *equation*, because it indicates the equality of the number and nature of the atoms before and after chemical action. On the left hand of the sign of equality are shown two molecules, and on the right hand two molecules; but, of the molecules on the left, one contains two atoms of hydrogen and the other

two atoms of chlorine, while of the molecules on the right each contains one atom of hydrogen and one of chlorine. The equation forms a short and convenient plan of recording the facts of experiment.

Instead of an equation, a *diagram* may be employed to exhibit the same fact. Thus :—



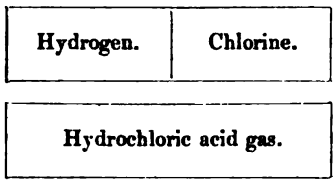

## PHYSICAL AND CHEMICAL CONSTITUTION OF MATTER.

### Relations of Gases, Liquids, and Solids.

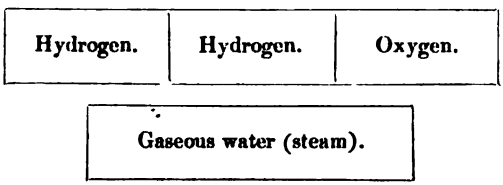
The molecules of a gas are not close together, for a quantity of gas may be compressed with very little force to half or one-fourth its bulk—in short, to such an extent that in many cases the molecules sufficiently approximate to form a liquid. In a liquid the molecules are still free to glide about with ease amongst each other; and though in solids they exhibit less mobility, still even solids may be compressed by powerful pressure, so that probably in no instance are molecules in absolute contact. (Moreover, from the researches of Caignard de la Tour, and, recently, of Andrews, there would seem to be no sharp lines of demarcation between the gaseous, liquid, and solid conditions of substances.) One's mental picture of the relative position of the molecules of gaseous, vaporous, liquid, or solid matter must be such a picture as that of the moving particles of dust in the air of a room, or such a relation to each other as that of the planets and stars suspended in space. There is abundant experimental evidence to warrant such a conception. A clear transparent fluid appears perfectly homogeneous, but is not so. Its particles are not in contact. Every one who has mixed 5 pints of rectified spirit with 3 pints of water knows that the 100 fluidounces of spirit and 60 fluidounces of water do not when mixed give 160 ounces of "proof" spirit, but only 156 ounces; the molecules of the liquids have gone closer together, having probably a little attraction for each other. Why a gas under pressure should immediately return to its original bulk when the pressure is removed, while a liquefied or solidified gas only slowly resumes the gaseous or vaporous state, is a question which requires for discussion a knowledge of the nature of forces other than the chemical. For it must be remembered that the study of the chemical force is mainly the study of the internal constitution of molecules, the study of the properties of entire molecules forming the domain of *Physics* —sometimes termed *Natural Philosophy*. (Physics, from φύσις, *phusis*, nature, that is, visible and material nature, the study of actions and reactions which do not involve entire and permanent

change in the properties of bodies—the study of the action of heat, light, electricity, magnetism, gravitation, etc., on matter.)

It is necessary, however, to state something more about the physical as well as the chemical conditions of the molecules of a *gas* in order that the learner may be prepared for the fact, that mixtures of certain gaseous elements, in combining to form gaseous compounds, diminish considerably in volume. Thus, while a pint of hydrogen and a pint of chlorine give a quart of hydrochloric acid gas,




two pints of hydrogen and one of oxygen are necessary to produce a quart of gaseous water (steam). It will be remembered that two volumes of hydrogen and one of oxygen were necessary in a previous experiment in which water was formed.




Now, that a pint of hydrogen gas and a pint of chlorine gas should, after chemical reaction or rearrangement of the atoms of the molecules has taken place, form two pints of hydrochloric acid gas, is quite what we should expect. For, first, the reader, by this time, is not astonished that the chemical combination is attended by entire change of properties; and, secondly, the experience of years has led him to expect that a pint of one thing added to a pint of another gives two pints of the mixture. But that two pints of hydrogen and one pint of oxygen should, after combination (and under like conditions of temperature and pressure), give, not three, but two pints of product (steam) is perhaps somewhat astonishing, and needs explanation. To this end let us picture a few of the molecules of hydrogen and as many molecules of chlorine. Draw with a pencil on paper several pairs of crosses (+ +) to represent hydrogen molecules and circles (° °) for chlorine molecules, or, if colored ink is at hand, red pairs of dots for hydrogen and green for chlorine. Or, at once, for facility in printing, let the following pairs of letters *h h* represent a few (say, nine) molecules of hydrogen, and *c c* molecules (nine) of chlorine—before combination.

| | | | | | |
|---|---|---|---|---|---|
| *h h* | *h h* | *h h* | *c c* | *c c* | *c c* |
| *h h* | *h h* | *h h* | *c c* | *c c* | *c c* |
| *h h* | *h h* | *h h* | *c c* | *c c* | *c c* |

Then, after combination, we shall have eighteen molecules of hydrochloric acid gas:—

| | | | | | |
|---|---|---|---|---|---|
| *h c* | *h c* | *h c* | *h c* | *h c* | *h c* |
| *h c* | *h c* | *h c* | *h c* | *h c* | *h c* |
| *h c* | *h c* | *h c* | *h c* | *h c* | *h c* |

But when two volumes of hydrogen and one of oxygen combine and give two volumes of steam, the mental picture must be not that of molecules somewhat nearer to each other than before, nor any difference in the size of the molecules, but a picture of molecules containing three instead of two atoms—thus, still using pairs of letters, just for the moment, to represent a few (the space will allow only twenty-seven) molecules:—

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| *h h* | *h h* | *h h* | *h h* | *h h* | *h h* | *o o* | *o o* | *o o* |
| *h h* | *h h* | *h h* | *h h* | *h h* | *h h* | *o o* | *o o* | *o o* |
| *h h* | *h h* | *h h* | *h h* | *h h* | *h h* | *o o* | *o o* | *o o* |

The twenty-seven molecules (eighteen hydrogen, nine oxygen) will, after combination, become eighteen molecules of steam:—

| | | | | | |
|---|---|---|---|---|---|
| *h o h* | *h o h* | *h o h* | *h o h* | *h o h* | *h o h* |
| *h o h* | *h o h* | *h o h* | *h o h* | *h o h* | *h o h* |
| *h o h* | *h o h* | *h o h* | *h o h* | *h o h* | *h o h* |

As already suggested, one's mental picture of a number of molecules may well give them such a relation to each other as that of a number of solar systems in the universe, equally distant from each other and each occupying a similar space, yet one system containing a sun and one planet, another a sun and two planets, and so on, or even one or more of the planets having one or more moons. Indeed the atoms in some very complex molecules really appear to have very much the relation to each other of the sun, planets, and moons of a solar system. To indicate such molecules by letters as above would, of course, require more space than is there given to the assumed pictures of molecules. For an amusing and philosophical picture of a complex molecule, the reader is referred to a pamphlet by Professor Clifford entitled "Atoms," by which he means physical atoms, physical units, or what chemists call molecules; it is published by F. Pitman, Paternoster Row, London, and its price is one penny.

Here occurs an opportunity that must not be lost of stating a mode of reasoning by which a molecule of oxygen (or of many other

elements) is shown to be a double structure—shown to contain two atoms. Five equal-sized bottles are before us, two filled with hydrogen, one with oxygen, and two with steam. (The bottles are hot enough to prevent the steam condensing to water, and all five are at the same temperature.) Apply heat so that all shall be equally heated, *the three different substances expand equally.* Cool equally, *the contents contract equally.* Apply equal pressure to all five, *each is equally affected.* Diminish pressure equally, *each portion of the three substances equally expands.* Gases (practically steam is a gas, it is simply not a permanent gas) thus similarly affected must be, physically, similarly constructed or constituted (a law which will again be referred to, on page 55); each bottle must contain the same number of particles or molecules, and at any one temperature and pressure the molecules in each must be equally distant from each other. We do not know what actual number or distance, but whatever be the number and distance it is the same for each bottle. Say that one million is the number, then we shall have a million of molecules in the first hydrogen bottle, a million in the second, a million in the oxygen bottle, and a million in each of the steam bottles. We will cause chemical combination between the two millions of hydrogen molecules and one million of oxygen molecules, producing (as we have seen) two millions of steam molecules, having the properties already stated. But a molecule of steam contains an atom of oxygen. Hence two millions of steam molecules contain two millions of oxygen atoms, which two millions of oxygen atoms have been obtained from one million of oxygen molecules. Therefore each molecule of oxygen was a double structure—each molecule of oxygen contained two atoms of oxygen. As Clifford says, "you cannot put 50 horses into 100 stables, so that there shall be exactly the same amount of horse in each stable; but you can divide 50 *pairs* of horses among 100 stables."

Thus much respecting the constitution of gaseous or vaporous matter. Our knowledge of the constitution of liquid and solid matter is still more limited.

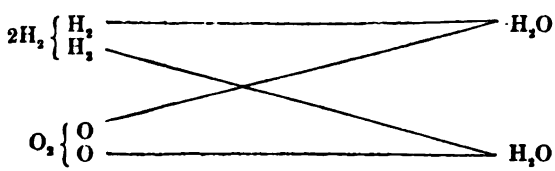
With regard to the notation of the subject, it will be sufficient to state here that a *symbol* usefully represents one volume of any gas, *a formula of any gas or vapor represents two volumes.* By remembering this general rule we may, by looking at a formula, tell how many volumes of constituents were concerned in the formation of a compound, and therefore what amount of condensation, if any, occurred during the act of formation. By thus reading and interpreting the formula for water, $H_2O$, we see that two volumes of steam (at any temperature) may be obtained from two volumes of hydrogen and one volume of oxygen (at the same temperature), and thus that the extent of condensation when hydrogen and oxygen (at a stated temperature) unite to form gaseous water (at the same temperature) is from three to two. This subject will again be treated of in connection with those of Chemical Combination and Specific Gravity of Gases.

## Further Remarks on General Chemical Notation.

We may now take an experiment already made as an additional example of chemical action, and describe the simplest way of expressing the same by notation. When two volumes of hydrogen and one of oxygen were caused to combine, the production of flame and noise proved that chemical action of some kind had taken place; had the experiment been performed in dry vessels, evidence of the precise action would have been found in the bedewment or moisture produced by the condensation of the water on the sides of the tube. Similar evidence was afforded on holding a cool glass surface over the hydrogen-flame. The action is expressed in the following equation:—

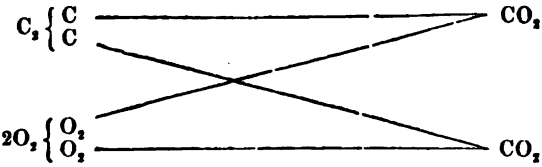
$$2H_2 + O_2 = 2H_2O.$$

Instead of an equation, the following diagram may be employed:—




The foregoing aggregation of symbols or short-hand characters, or *formula*, $H_2O$, is, then, a convenient picture of the facts that have already come before us, viz., that water is formed of the elements hydrogen, H, and oxygen, O; moreover, that it is formed of two measures or volumes of hydrogen, $H_2$, to one of oxygen, O; and, thirdly, that the molecule of water ($H_2O$) is formed of two atoms of hydrogen ($H_2$) and one of oxygen (O). The formula also fulfils the fourth function of indicating that the two volumes of hydrogen and one of oxygen in combining condensed to two volumes of steam. That the resulting bulk of steam afterwards shrunk most considerably in condensing to water is another matter altogether, a physical and not a chemical result, and due to the approximation of the molecules of water *after* formation.

Another experiment already performed, illustrating the character of the manifestations of chemical force (symbolically noted as follows), was that in which the red-hot carbon of wood was plunged into oxygen. The evidence of chemical action in that case was the sudden inflammation of the carbonaceous extremity of the wood. The particles of carbon and oxygen, having intense attraction or affinity for each other at that temperature, rushed together so impetuously as suddenly to produce a large additional quantity of heat, an amount sufficient to cause the particles to emit an intense white light. The action between carbon and oxygen is expressed on paper in either of the following ways:—




$CO_2$ is the formula of the well-known gaseous body commonly termed carbonic acid gas.

The reader should here draw for himself equations or diagrams similar to those on pages 46 and 47, and thus show the formation of the three bodies he has already produced—namely, phosphoric anhydride ($P_2O_5$), sulphurous acid gas ($SO_2$), and iodide of iron ($FeI_2$), submitting the same, if possible, to a tutor or other authority to assure himself of their correctness.

*Note.*—In the foregoing experiments several illustrations occur of the formation of compounds having the gaseous, liquid, and solid conditions, in one of which three forms all matter in the universe exists.

### Laws of Chemical Combination (by Weight).

Chemistry as a science is little more than a hundred years old, though very many of the facts and operations we now term chemical have been known as isolated items of knowledge for centuries. Thus, the ancient Egyptians made glass, vitriol, soap, and vinegar; and the Greeks started the idea that matter was composed of a few elements, imagining earth, air, fire, and water to be elements. But the great general principles which interlace and bind together separate facts, those which from their extensive application and importance are denominated *laws*, have all been brought to light since the year 1770.

### *First Law relating to Chemical Combinations.*

Between 1785 and 1800, Bryan Higgins, William Higgins, Wenzel Richter, and Proust made analyses and researches which led up to the following generalizations: *When compounds unite to form definite chemical substances, they always combine in the same proportions.* The curious character of this fact could but be most striking, and indeed is so now, to the mind receiving it for the first time. Thus water (a compound) added to quicklime (a compound) gives slaked lime, a perfectly definite chemical substance. But whereas sand and water, sugar and water, sand and sugar, and such *mixtures* may be obtained by adding together the ingredients in any proportions whatever, say, 90 of sugar and 10 of sand, or 10 of sugar and 90 of sand, slaked lime (say 100 parts) invariably results from the combination of 75¾ of quicklime and 24¼ of water. If a larger proportion than 75¾ per cent. of quicklime be employed, the excess remains as quicklime mixed with the slaked lime; and if more than 24¼ per cent. of water be used, an excess of water remains with the slaked lime and

evaporates if the mixture be exposed to the air. Dalton discovered that when *elements* unite to form a definite substance, they, like compounds, always combine in the same proportions; and he was the first to set forth the law in a manner which was at once clear and comprehensive enough to include the former generalization. Thus:—

*A definite compound always contains the same elements in the same proportions.*

Take another example. Common salt always contains 39⅓ per cent. of the metal sodium to 60⅔ of chlorine, and water always 89 of oxygen to 11 per cent. of hydrogen (more exactly 88.89 to 11.11). As with quicklime and water, so with the chlorine and sodium, and the constituents of many (not all) chemical compounds; in such cases if either be added to the other in any quantity beyond stated proportions, the excess plays no part whatever in the act of combination. (In *some* cases, as will be seen directly, excess of either plays a very simple but very remarkable part.) In short, whether a compound be made directly from its elements, or by the combination of other compounds, or indirectly as one of two products of the action of substances chemically on each other, whatever be its origin, if it is a definite compound it always contains the same elements in the same proportions.

### *Second Law relating to Chemical Combinations.*

Dalton further made such experimental researches as enabled him to lay down a second great law. He found that, while many substances only united chemically in one proportion, others combined in two or even more, and he studied several such naturally related bodies. He found that while carbonic oxide (a gas formed when charcoal is burned with an insufficient supply of air) contains such a proportional weight of carbon and oxygen as is represented by (to use the simplest figures) 3 and 4, carbonic acid (a gas formed when charcoal is burned with excess of air) contains 3 of carbon to exactly twice 4 of oxygen. He proved that a similar relation existed between two compounds of carbon and hydrogen, and between a cluster of compounds of nitrogen and oxygen. The first of the latter, to a given quantity of nitrogen, contains a certain proportion of oxygen; the next, to the same quantity of nitrogen, had twice the proportion of oxygen: and the others have respectively three, four, and five times as much oxygen as the first, the quantity of nitrogen remaining the same throughout. Dalton thus generalized these facts:—

*When two elements unite in more than one proportion, the resulting compounds contain, to a constant proportion of one element, simple multiple proportions of the other—or the weights of the constituent elements bear some similar simple relation to each other.*

Thus carbonic oxide gas is a definite compound always containing fixed proportions of carbon and oxygen, and carbonic acid gas also a

definite compound always containing fixed proportions of carbon and oxygen. Both thus obey the first law of combination. But whereas carbonic oxide contains, or may be made from, 30 parts (ounces, grains, or other weights) of carbon and 40 of oxygen, carbonic acid contains, or may be made from, 30 parts of carbon and exactly twice 40 of oxygen.

This second law cannot but be as striking as the first when freshly unveiled to the mind. Sand and sugar, or any substances which do not act *chemically* on each other, may be mixed in the proportions of 30 to 40, 30 to 80, 30 to 60, or any other quantities; but if an attempt be made to burn 30 parts of carbon in 60 of oxygen, the elements will themselves naturally assert their own special combining powers, and refuse, so to say, to unite in these proportions: the 30 of carbon will first combine with 40 of oxygen and form 70 of carbonic oxide, and this gas, which, had it the opportunity, would combine with 40 more of oxygen and form carbonic acid gas, finding only half that quantity, namely, 20 of oxygen present, contents itself by one-half (that is, 35 of carbonic oxide) accepting the 20 of oxygen and becoming carbonic acid gas, while the other half remains as carbonic oxide. This is a most wonderful fact. Again, if 30 parts of carbon be burnt in more than 80, say 85, of oxygen, only 80 will be used, the other 5 remaining as oxygen merely *mixed* with the resulting carbonic acid gas. If we attempt to burn 30 parts of carbon in less than 40 of oxygen, the oxygen will take up three-fourths its weight of carbon and form carbonic oxide, while the excess of carbon will remain as carbon.

### Recapitulation.

Nature does not always permit man to mix things in any proportions he pleases. She does sometimes. She does if he only stirs things together, or if he only uses the attractions of adhesion or cohesion in binding the materials together; but if he employs chemical attraction, she restricts him to special proportions. That is to say, if the things mixed do not attack one another or intimately combine, then admixture may be effected in any proportion; and the mixture is a mere mixture having the mean properties of its components. Examples of such mixtures are seen in compound plasters, pill-masses, confections, and plum-puddings. But if the things do unite to form, not a mere mixture having the mean properties of its components, but a compound having new and distinct and definite characters of its own, then nature does not permit man to mix the things in any proportion he pleases. The proportion is one fixed and constant; and if he substitutes proportions of his own, the things unite in the proportions fixed by nature, and the excess he has added either remains in its original uncombined condition, or it combines with the compound already produced to form a second *different* compound. Any one compound, that is, the *same* compound, always contains the same elements in the same proportions, and can only be made from the same elements in the same proportions. An attempt to mix the same elements in other proportions would result in one of two failures, namely, either the extra propor-

tion would remain free and uncombined, or it would combine and convert the first compound, or a portion of it, into a *different* compound. The fresh compound thus produced, like the first, and indeed like all definite compounds, of course always contains the same elements in the same proportions.

In short (law 1) any definite compound always contains the same elements in the same proportions, and (law 2) any two elements uniting in more than one proportion unite in multiplies of that proportion, and produce so many different definite compounds. Taking hydrogen as uniting in proportions of 1, oxygen unites in proportions of 16—that is, 16, twice 16, thrice 16, and so on, never in intermediate proportions. Carbon unites in proportions of 12, sulphur of 32, chlorine of 35½. And every other element has its combining proportion fixed by nature.

The student of chemistry is recommended to accept these two great natural facts, great enough to be dignified by the name of *laws*, in all their inherent solidity and simplicity. Of course, he will wonder *why* substances should combine, chemically, only in fixed proportions when forming a definite body, and *why*, when a substance combines in more than one proportion to form different compound bodies, the proportions should only be multiple proportions; and an extremely ingenious and useful explanation has been suggested by Dalton (see the following paragraphs on the theory that matter is built up of atoms); but man has not yet succeeded in so questioning nature as to gain from her a *satisfactory* answer to such questions; hence, until he does succeed, any hypothesis, such as Dalton's, should be held intelligently but loosely. The facts themselves, however, should be grasped with the student's utmost tenacity.

Careful consideration of the foregoing two great laws has suggested an important truth sometimes termed *The Law of Chemical Combination*, namely: *The proportions in which two elements unite with a third are the proportions (or simple multiples or submultiples of the proportions) in which they unite with each other.* Thus oxygen in proportions of 16 unites with hydrogen, and carbon in proportions of 12 unites with hydrogen; therefore 16 and 12 are the proportions in which oxygen and carbon will unite with each other.*

### The Atomic Theory.

The laws which Dalton (1803 to 1808) so largely aided to unveil—two grand and wonderful truths—he explained and correlated by a simple and beautiful hypothesis. *Dalton suggested that matter was*

* See Axiom 1 in Hawtrey's fascinating "Introduction to the Elements of Euclid," Longmans & Co., London, 2*s.* 6*d.*, a book strongly recommended to any chemical student who is not familiar with the mode of reasoning commonly termed geometrical.

*not infinitely divisible, but composed of minute particles or* ATOMS *having an invariable character.* In the words of Wurtz, "*To an old and vague notion he attached an exact meaning by supposing that the atoms of each kind of matter possess a constant weight, and that combination between two kinds of matter takes place not by penetration of their substance, but by juxtaposition of their atoms.*"

Thus under this hypothesis, or atomic theory as it is generally termed, carbonic oxide is a definite compound always containing the same elements in the same proportions, *because* each particle of it is composed of an atom of carbon and an atom of oxygen chemically united, the weights of the atoms being in the proportion of 3 and 4, that is, having a constant weight of 12 and 16, as we now believe. Carbonic acid gas is also a definite compound always containing the same elements in the same proportions, and the proportion of oxygen is just double that in carbonic oxide, *because* each particle of it is composed of an atom of carbon (weighing 12) and two atoms of oxygen (each weighing 16).

Imaginary pictures of molecules of carbonic oxide gas and carbonic acid gas.*

Again, the facts that with 12 of carbon oxygen unites in the proportion of 16, or a multiple of 16; that with 12 of carbon sulphur unites in the proportion of 32, or a multiple of 32 (the liquid known as disulphide of carbon is a chemical compound of 12 of carbon to twice 32 of sulphur); and, thirdly, that oxygen and sulphur unite in proportions of 16 and 32, are at once explained on the assumption that these elements exist in atoms which have the respective weights mentioned. Existing in indivisible particles (atoms) which weigh 16, 12, and 32, oxygen, carbon, and sulphur *must* unite in indivisible weights of 16, 12, and 32.

### ATOMIC WEIGHTS.

What has just been stated respecting two or three elements is true of all the elements. It is a fact, that, when elements unite with one another in the peculiar and intimate manner termed chemical, they do not combine in the haphazard proportions of a mere mixture, but in one fixed and constant proportion. Such proportions or weights represent, according to Dalton, the weights of their atoms. Oxygen

* The size of atoms, their shape, their absolute weight—whether or not they are in actual contact—whether or not they are fixed in relation to each other, free to move about each other, or in a constant state of motion—and whether or not the chemical force actuates them as the force of gravitation influences our earth, and moon, and solar systems, are matters of which at present we know almost nothing. The two pictures are not intended to convey any impression that the following formulæ do not give: CO or OC, OCO or OOC or COO or $CO_2$.

unites with other elements in proportions of 16, therefore 16 is the weight of the atom of oxygen. Chlorine unites with other elements in proportions of 35½, therefore 35½ is the atomic weight of chlorine. And for a similar reason the atomic weights of hydrogen will be 1, carbon 12, sulphur 32, nitrogen 14, and iodine 127. Of course it will be understood that these are the *relative* weights of atoms, for we cannot know the absolute weights. All that is known is that the chlorine atom, for instance, is 35.5 times as heavy as the hydrogen atom, whatever the absolute weight of the latter may be, and the iodine atom 127 times as heavy. The quantity of metal which with 35.5 of chlorine will form a chloride, and with twice 35.5 a second chloride (dichloride or bichloride) will require 127 of iodine to form an iodide, and twice 127 of iodine to form a second iodide (a diniodide or biniodide).*

*Note on Notation.*—A fourth function of a symbol is to represent atomic weight. Thus the symbols H, Cl, O, etc., not only perform the office of representing (*a*) names, (*b*) single volumes, and (*c*) single atoms, but (*d*) definite weights of the respective elements.

$$H = 1,\ Cl = 35.5,\ O = 16,\ I = 127,\ N = 14,\ K = 39,\ \text{etc.}$$

### Laws of Chemical Combination (by Volume).

In 1809 Gay-Lussac showed it to be a fact that when gaseous elements unite with one another in the intimate manner termed chemical, they do not combine in the haphazard proportions (that is, proportions by measure or volume) of a mere mixture, but in constant proportions in the case of any single definite compound, and in simple multiple proportions in cases where two elements form more than one definite compound. He thus proved that the laws respecting the constancy of weight with which elements combine hold good with reference to volume, at all events in those cases in which elements exist in or can be made to assume the gaseous condition. A volume of hydrogen gas and an equal one of chlorine gas give hydrochloric acid gas. Two volumes of hydrogen and one of oxygen give water-vapor or steam. Such volumes or simple multiples are alone the proportions by bulk in which elements combine. If any excess of either gas be mixed and combination attempted, only the stated proportions really combine, the excess remaining unaltered. Further, following Gay-Lussac, on weighing these similar and equal volumes of hydrogen, chlorine, and oxygen, we find that the chlorine is 35.5 times as heavy as hydrogen, and oxygen 16 times as heavy as hydrogen.

In 1811 and 1814, Avogadro and Ampère, reasoning on the fact that all gases are similarly affected by variations of pressure (Boyle, 1662, verified by Mariotte) and temperature (Charles), concluded that all gases must be similarly constituted—similarity in properties

* Only the atomic weights of the above and a few of the chief metallic elements need be committed to memory; others can be sought out as occasion may require. A complete Table of Atomic Weights is given at the end of the volume.

always indicating similarity in character or nature; in other words, that, if *equal volumes of gases* be taken *under like conditions*, each will *contain the same number of molecules*, similar in *size* and equally distant apart. The deduction is obvious. *The weights of molecules of gaseous elements (that is, pairs of atoms, and therefore of atoms themselves) must differ to the extent that the weights of equal volumes of those elements differ.* Equal volumes of hydrogen, chlorine, and oxygen, weighing respectively 1, 35.5, and 16, and each of these volumes containing an equal number of molecules, each formed of two atoms, it follows that the relative weights of the atoms will be 1, 35.5, and 16.

It will thus be seen that the weight of the *volume* in which an element combines, and the *actual weight* in which it combines, irrespective of volume, are identical. For instance, we should find by experiment that, as a simple matter of fact, oxygen unites with other elements in proportions of 16 by weight, while hydrogen combines in proportions of 1. Turning, then, to experiments on the volumes in which hydrogen or oxygen combine, and having ascertained those volumes, and then having weighed them, we should find that the oxgyen volume weighs 16. while the hydrogen weighs 1. In compounds in which hydrogen was found in proportions of 1 grain, oxygen would be found in proportions of 16 grains. In gaseous compounds in which hydrogen was found in proportions of, say, 27 ounces by measure, oxygen would be found in proportions of 27 ounces by measure; the 27 ounces of hydrogen would be found to weigh 1 grain, and the 27 ounces of oxygen to weigh 16 grains.

Thus the two great facts or laws respecting chemical compounds which Dalton laid down by ascertaining the exact weights in which bodies combine, Gay-Lussac confirmed by experiments on the exact volumes in which elements combine. Further, Gay-Lussac's experiments and Avogadro's reasoning strongly supported Dalton's theory of atoms.

## Recapitulation.

What are atomic weights or combining weights? First, they are represented by the smallest proportion (relative to 1 part of hydrogen) in which an element migrates from compound to compound. Thus 1 part by weight of hydrogen can be eliminated from 18 similar parts of water by action of certain metals, leaving 1 of hydrogen and 16 of oxygen combined with the metal. From the latter compound 1 more of hydrogen is eliminated by a second experiment with more metal, leaving 16 of oxygen combined with the metal. In these and other well-known reactions 16 parts of oxygen take part in the various operations; 16, therefore, is the probable atomic weight of oxygen; and so with other elements and radicals. Secondly, the weights of the atoms, or the atomic weights, of the gaseous elements already studied, must differ from each other to the extent that equal volumes of those elements differ in weight. For equal volumes of an element contain an equal number of molecules equal in size (Avogadro's and Ampère's law), and each molecule is composed of two atoms; so that equal volumes of elements contain an equal number

of atoms. Now, bulk for bulk, chlorine is thirty-five and a half (35.5) times as heavy as hydrogen; so that the molecule of chlorine must be 35.5 times the weight of the molecule of hydrogen; for molecules are equal in bulk. And as the molecules of chlorine and hydrogen contain two atoms each, the atom of chlorine must be 35.5 times as heavy as that of hydrogen. The actual weight of atoms can never be ascertained, but that is of little consequence if we can only determine, with exactitude, their comparative weights. Comparing, then, all atomic weights, sometimes obscurely termed equivalents, with each other, and selecting hydrogen as the standard of comparison (because it is the lightest body known, and therefore, probably, will have the smallest atomic weight), and assigning to it the number 1, we see that the atomic weight of chlorine will be represented by the number 35.5. By parity of reasoning the atomic weight of oxygen is 16; for oxygen is found, by experiment, to be 16 times as heavy as hydrogen. Similarly the atomic weight of nitrogen is found to be 14. The atomic weight of carbon is 12—not because its vapor has been proved to be 12 times as heavy as hydrogen, for it has never yet been converted into the gaseous state, but because no gaseous compound of carbon, which has been analyzed, has been found to contain in 2 volumes (one of which, if hydrogen, would weigh 1 part) less than 12 parts of carbon.

By thus weighing equal volumes of gaseous elements, or equal volumes of gaseous compounds of non-volatile elements, and ascertaining by analysis the proportion of the non-volatile element whose atomic weight is being sought to the volatile element whose atomic weight is known, the atomic weights of a large number of the elements have been determined. Some of the elements, however, do not form volatile compounds of any kind; the stated atomic weights of these elements, therefore, are at present simply the proportions by weight in which they combine with or displace elements whose atomic weights have been determined, the proportion being in most cases checked by isomorphic considerations and the relation of the element to other forces, especially heat.* (*Vide infra.*)

### Molecular Weight and Molecular Volume.

*The weight of the molecule* is simply the sum of the weights of its atoms; thus

$$H_2 = 2,\ O_2 = 32,\ Cl_2 = 71,\ H_2O = 18,\ HCl = 36.5.$$

* *Isomorphous* bodies (from ἴσος, *isos*, equal, and μορφή, *morphē*, form) are those which are similar in the shape of their crystals. The identity in crystalline form is so commonly associated with similarity of constitution that non-crystalline substances resembling each other in structure are often regarded as isomorphous. When one element unites with another in more than one proportion, and its atomic weight is so far uncertain, the isomorphism of either of its compounds with some other compound of known constitution is usually accepted as decisive evidence as to which proportion is atomic. The specific heat of elements will be treated of subsequently.

*Molecular Volume.*—If the quantities just mentioned be weighed out (in grains or other weights), or if the molecular weight of any gases or liquids be taken and exposed to similar (high) temperatures and pressures, *they will all be found to occupy the same volume.* Conversely, if equal volumes of gases or vapors be measured out, and then the whole weighed, the resulting figures (all referred to 2 of hydrogen as a starting-point or standard) are the molecular weights of the respective substances. Thus a volume of hydrogen (about half a gallon) which, at a temperature of, say, 300° F. or 400° F., and common atmospheric pressure, would weigh 2 grains, would in the case of vapor of water (steam) weigh 18 grains. Hence we are justified in considering, indeed compelled to consider, the molecule of water to contain two atoms of hydrogen (= 2) and one of oxygen (= 16), and its formula to be $H_2O$ (= 18), and not $H_4O_2$, in which case its vapor would be twice as heavy as it really is found to be.

*Construction of Formulæ.*—The composition of hydrochloric acid (HCl), water ($H_2O$), ammonia ($NH_3$), carbonic acid gas ($CO_2$), or any other compound, as well as the weight of an element that may be concerned in its formation, cannot be ascertained by actual experiment until the student is far advanced in practical chemistry—until he is able to *analyze* not only *qualitatively*, but, by help of a balance, *quantitatively*. The percentage composition of a substance having been determined by quantitative analysis, its formula is constructed by aid of the foregoing and other theoretical considerations. The correctness of such formulæ can be verified by expert analysts, but must be taken for granted by learners. This subject will again be referred to in the latter part of this Manual.

## Quantivalence of Atoms.

Turning from the *weights* of atoms, their *value* may now be considered; their *quantivalence* (from *quantitas*, quantity, and *valens*, being worth) may be stated. The chemical value of atoms in relation to each other may be compared to the exchangeable value of coins. As compared with a penny (1*d.*) a groat (4*d.*) is four-valued; as compared with hydrogen carbon is quadrivalent. Here again hydrogen is conveniently adopted as the standard of comparison. An atom of oxygen in its relations to an atom of hydrogen is bivalent (pronounced thus, biv'-a-lent; of double worth, from *bis*, twice, and *valens*); an atom of it will displace two atoms of hydrogen, or combine with the same number; nitrogen is usually trivalent (triv'-a-lent; from *tres*, three, and *valens*); and carbon, quad-riv'-a-lent (from *quatuor*, four, or *quater*, four times, and *valens*). Chlorine, iodine, and bromine, as well as potassium, sodium, and silver among the metals, are, like hydrogen, univalent (u-niv'-a-lent; from *unus*, one, and *valens*). Barium, strontium, calcium, magnesium, zinc, cadmium, mercury, and copper, like oxygen, are bivalent. Phosphorus, arsenicum, antimony, and bismuth, like nitrogen, usually exhibit trivalent properties; but the composition of certain compounds of these five elements shows that the several atoms are sometimes quinquivalent (quin-quiv'-a-lent; *quinquies*, five times, and

*valens*). Gold and boron are trivalent. Silicon (the characteristic element of flint and sand), tin, aluminium, platinum, and lead resemble carbon in being quadrivalent. Sulphur, chromium, manganese, iron, cobalt, and nickel are sexivalent (sex-iv'-a-lent; from *sex*, six, or *sexies*, six times, and *valens*), but frequently exert only bivalent, trivalent, or quadrivalent activity. This *quantivalence* (quant-iv'-a-lence; from *quantitas*, quantity, and *valens*), also termed *atomicity* (maximum quantivalence), *dynamicity*, and *equivalence* of elements, may be ascertained at any time on referring to the Table of the Elements at the end of this volume, where Roman numerals, I, II, III, IV, V, VI, are attached to the symbols of each element to indicate atomic univalence, bivalence, trivalence, quadrivalence, quinquivalence, or sexivalence. Dashes (H′, O″, N‴) similar to those used in accentuating words are often used instead of figures in expressing quantivalence. The quantivalence of elements, as they one after another come under notice, should be carefully committed to memory; for the composition of compounds can often be thereby predicated with accuracy and remembered with ease. For instance, the hydrogen compounds of chlorine, Cl′, oxygen, O″, nitrogen, N‴, and carbon, C⁗, will be respectively H′Cl′, $H'_2O''$, $H'_3N'''$, and $H'_4C''''$,—one univalent atom, H′, balancing or saturating one univalent atom Cl′; two univalent atoms $H'_2$, and one bivalent atom O″, saturating each other; three univalent atoms, $H'_3$, and one atom having trivalent activity, N‴, saturating each other; and four univalent atoms, $H'_4$, and one quadrivalent atom, C⁗, saturating each other. Carbonic acid gas, $C^{IV}O^{II}_2$, again, is a saturated molecule containing one quadrivalent and two bivalent atoms.

*The subject of quantivalence will be further explained after the first six metals have been studied, when abundant illustrations of it will have occurred.*

---

## DEFINITIONS.

*Chemistry* is the study of the chemical force.

*The Chemical Force*, like other forces, cannot be described, for, like them, it is only known by its effects. It is distinguished from other forces by the facts that (*a*) it produces an entire change of properties in the bodies on which it is exerted, and (*b*) that it is exerted only between definite weights and volumes of matter. Like the *force of cohesion*, which is the name given to the attraction which molecules have for each other, and which is great in solids, small in liquids, and apparently absent in gases, and like the *force of adhesion*, which is the name given to the attraction which a mass of molecules has for another mass, *the chemical force* acts only within immeasurable distances: indeed, inasmuch as the chemical force appears to reside in atoms, that is to say, is exerted inside a molecule, while all other forces affect entire molecules, the chemical force may be said to be distinguished (*c*) by being exerted within a smaller distance than that at which any other force is exerted.

*An Element* is a substance which cannot by any known means be resolved into any simpler form of matter.

*An Atom* of any element is a particle so small that it undergoes no further subdivision in chemical transformations.

*A Molecule* is the smallest particle of matter that can exist in a free state.

*A mere Mixture* of substances is one in which each ingredient retains its properties.

*A Chemical Compound* is one in which definite weights of constituents have combined, and during combination have undergone an entire change of properties. A "compound" in pharmacy is an intimate mixture of substances, but still only a mixture; it is not a *chemical* compound, the ingredients have not entered into *chemical* union or combination.

*Combustion* is a variety of chemical combination, a variety in which the chemical union is sufficiently intense to produce heat and, generally, light.

*The Law of Diffusion* is one under which gases mix with each other at a rate which is in inverse proportion to the square root of their relative weights; that is, irrespective of and even in spite of their comparative lightness or heaviness.

*A Chemical Symbol* is a capital letter, or a capital and one small letter. It has four functions, namely:—

1. It is short-hand for the name of the element.
2. It represents one atom of the element.
3. It stands for a constant weight of the element—the atomic weight or combining weight.
4. Symbols represent single and equal volumes of gaseous elements.

*A Chemical Formula* represents a molecule either of an element or of a compound. It also has four functions:—

1. It indicates at a glance the names of the elements in the molecule.
2. Its symbol or symbols, together with a small figure attached to the foot of any symbol, show the number of atoms in the molecule.
3. It stands for a constant weight of a compound—the molecular weight, that is, the sum of the weights of the atoms in the molecule.
4. It represents two volumes of the substance in the state of gas or vapor. (In the case of bodies which cannot be volatilized this statement is only probably correct.)

*A Chemical Equation* or *a Chemical Diagram* is a collection of formulæ and symbols so placed on paper as to form a picture or illustration of the state of things before and after that metathesis (interchange of atoms) of molecules which results in the formation of molecules of new substances.

*A Solid* is a substance the molecules of which are more or less immobile, though probably not in absolute contact.

*A Liquid* is a substance the molecules of which so freely move about each other that it readily assumes and retains the form of any vessel in which it is placed.

*A Gas* is a substance the molecules of which are so far apart that they seem to have lost all attraction for each other, and, indeed, to have acquired the property of repulsion to such an extent that they are only prevented from receding to a still greater extent by the pressure of surrounding matter. Motion is especially characteristic of gaseous fluids.

*The Three Laws regulating Chemical Combination (either by weight or volume).*

*First.* A definite compound always contains the same elements and the same proportions of those elements—by weight or volume.

*Second.* When two elements unite in more than one proportion, they do so in simple multiples of that proportion.

*Third.* The proportions in which two elements unite with a third are the proportions in which they unite with each other.

*Atomic Weights* are, first, the proportions in which elements are found to combine with each other by weight. (The figures showing these proportions are purely relative, but all chemists agree to make this relation fixed by giving the number 1 to hydrogen.) Secondly, they are the weights of equal volumes of elements in the state of gas (relative to 1 of hydrogen).

*Molecular Weights.*—These are the weights of equal volumes of gases or vapors, under equal circumstances of temperature and pressure, and relative not to 1 but to 2 of hydrogen. In the case of non-volatile bodies, molecular weight is deduced from the observed analogies of the bodies with those whose molecular weight admits of proof.

*Quantivalence of Atoms.*—The observed power, force, or value for work of an atom—relative to 1 of hydrogen.

---

THE LEARNER IS RECOMMENDED TO READ THE FOREGOING PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY CAREFULLY ONCE OR TWICE, THEN TO STUDY (EXPERIMENTALLY, IF POSSIBLE) THE FOLLOWING PAGES, RETURNING TO AND READING OVER THE GENERAL PRINCIPLES FROM TIME TO TIME UNTIL THEY ARE THOROUGHLY COMPREHENDED.

Minor Principles and Generalizations will be found scattered throughout the following pages.

Students of pure chemistry, especially when fairly well acquainted with chemical facts, will find the Principles of Chemistry, including the probable Constitution as distinguished from the mere Composition of Chemical Substances, amply set forth in Tilden's "Chemical Philosophy," one of Longman's series of "Text-books of Science."

## QUESTIONS AND EXERCISES.

42. What do you understand by *chemical* action? Give examples.

43. How is the chemical force distinguished from other forces?

44. Adduce evidence that elements exist in compounds; that sulphide of iron, for instance, still contains particles of sulphur and iron, though it possesses properties so different from these elements.

45. Define the term atom.

46. What condition is essential for the manifestations of chemical force?

47. Can an atom exist in an uncombined state? and when are the atoms of an element most potent to enter into chemical combination?

48. What is a molecule.

49. How may the results of chemical reactions be expressed on paper?

50. Enumerate the functions of a symbol.

51. Give the additional functions of a chemical formula.

52. Describe by a diagram or an equation the reaction which ensues when red-hot charcoal is plunged into oxygen gas.

53. Draw diagrams representing the formation of $P_2O_5$, $SO_2$, and $FeI_2$ respectively.

54. Enumerate the differences between the physical state of the molecules in a solid, a liquid, and a gas.

55. State the law of constant proportions.

56. State the law of multiple proportions.

57. State the law of reciprocal proportions.

58. Give illustrations of the above laws.

59. Describe the origin and use of the atomic theory.

60. What do you understand by the atomic weight, and the molecular weight of an element?

61. Representing the weight of an atom of hydrogen as 1, what will be the atomic weights of carbon, sulphur, nitrogen, and iodine? Give reasons for considering the stated weights to be correct.

62. In what proportion, by volume, do elements in the gaseous state chemically combine?

63. What relation exists between the volumes of elements in the gaseous state and their atomic weights? Give the explanation for this.

64. Is there any difference between the molecular volume of a simple or of a compound gas?

65. Define isomorphism.

66. Explain the value of isomorphism as evidence of atomic weight.

67. What is to be understood by the quantivalence of an element? Give examples of univalent, bivalent, trivalent, and quadrivalent atoms.

68. How may the quantivalence of an element be expressed in its atomic symbol?

69. Give the formulæ of two or three compounds in which the

quantivalence of one atom is saturated by the combined quantivalence of others.

The reader is also recommended to question himself, or be questioned, on the "definitions" given on pages 56, 57, and 58.

---

## THE ELEMENTS AND THEIR COMPOUNDS.

Having thus obtained a general idea of the nature of such elements as have especial interest for the medical and pharmaceutical student, and which indeed are all with which any student of chemistry should at present occupy his attention, we may pass on to consider in detail the relations of the elements to each other. The elements themselves, in the free condition, are seldom used in medicine, being nearly always associated, bound together by the chemical force; in this combined condition, therefore, they must be studied. Each combination of elements or chemical compound will, in the following pages, be regarded as containing two parts or roots, two radicals; the one usually metallic, or, to speak more generally, basylous; the other commonly a non-metallic, simple or complex, acidulous radical. The basylous radicals, or metals, will be considered first, the acidulous radicals afterwards. Each radical will be studied from two points of view, the synthetical and the analytical: that is to say, the properties of an element on which the preparation of its compounds depends will be illustrated by descriptions of actual experiments, and thus the principles of chemistry, and their application to medicine and pharmacy, be simultaneously learnt; then the reactions by which the element is detected, though combined with other substances, will be performed, and so the student will be instructed in qualitative analysis. Synthetical and analytical reactions are, in truth, frequently identical, the object with which they are performed giving them synthetical interest on the one hand, or analytical interest on the other.

A good knowledge of chemistry may be acquired synthetically by preparing considerable quantities of the salts of the different metals, or analytically by going through a course of pure qualitative analysis. But the former plan demands a larger expenditure of time than most students have to spare, while under the latter system they generally lose sight of the synthetical interest which attaches to analytical reactions. Hence the more useful system, now offered, of studying each metal, etc., from both points of view, time being economized by the operator preparing only small specimens of compounds.

Chemical synthesis and analysis, thoughtfully and conscientiously followed, will insensibly carry the principles of chemistry into the mind, and fix them there indelibly.

## POTASSIUM.

Symbol K. Atomic weight 39.

Formula $K_2$. Probable molecular weight 78.

*Memoranda.*—The chief sources of the potassium salts* are the chloride found at Stassfurt, in Prussia, in the form of the mineral *Carnallite* (chloride of potassium 50, chloride of sodium 25, and chloride of magnesium 25, in 100 parts); the nitrate found in soils, especially in warm countries; and the compounds of potassium existing in plants. *Kainit*, a double sulphate of potassium and magnesium, also occurs among the Stassfurt minerals. The vegetable salts of potassium are converted into carbonate (other salts are present) when the wood or other parts are burned to ashes. If the ashes be lixiviated with water, and the solution evaporated to dryness, the residue, when fused, constitutes *crude potashes*. The residue, calcined on the hearth of a reverberatory furnace till white, gives the product termed *pearlash* (*Potassii Carbonas Impura*, U. S. P.). Large quantities of carbonate are thus produced in North America and Russia, and, latterly, from the sugar beet marc, in France. From the native chloride and from the carbonate purified by treating the pearlash with its own weight of distilled water, filtering and evaporating the solution until it thickens, and stirring constantly, "so as to form a granular salt" (*Potassæ Carbonas*, B. P.; *Potassii Carbonas*, U. S. P.), nearly all other compounds of potassium are made. Exceptions occur in cream of tartar (*Potassæ Tartras Acida*, B. P.; *Potassii Bitartras*, U. S. P.), which is simply the purified natural potassium salt of the grape-vine, and in nitrate of potassium. Potassium is a constituent of between forty and fifty chemical or Galenical preparations of the British Pharmacopœia.

Carbonate of potassium is a white crystalline or granular powder, insoluble in alcohol, very soluble in water, rapidly liquefying in the air through absorption of moisture, alkaline and caustic to the taste. It loses all water at a red heat. *Potassii Carbonas Pura*, U. S. P., is obtained by heating the bicarbonate to redness: the resulting white anhydrous carbonate is converted into hydrous granular carbonate by solution in water and evaporation until a dry granular salt remains.

*Preparation.*—Potassium itself is isolated with some difficulty by distilling a mixture of its carbonate and charcoal. It rapidly oxidizes in the air, and hence is always kept below the surface of mineral naphtha, a liquid containing no oxygen.

*Quantivalence.*—The atom of potassium is univalent, K'.

---

* The ill-defined term *salt* includes most solid chemical substances, but more especially those which assume a crystalline form.

REACTIONS HAVING (*a*) SYNTHETICAL AND (*b*) ANALYTICAL INTEREST.

(*a*) *Synthetical Reactions.*

These are actions utilized in manufacturing preparations of potassium. The word *synthesis* is from σύνθεσις (*sŭnthĕsis*), a putting together, as opposed to analysis, from ἀναλύω (*analuo*), I resolve.

## Hydrate of Potassium. Caustic Potash.

*First Synthetical Reaction.*—Boil together, for a few minutes, in a test tube, five or six grains of carbonate of potassium ($K_2CO_3$) and a like quantity of slaked lime (Ca2HO) with a small quantity of water. Set the mixture aside in the test-tube rack till all solid matter has subsided.

This liquid is a solution of caustic potash, or hydrate of potassium (KHO). Made of a prescribed strength, it forms the *Liquor Potassæ*, B. P. (5.84 per cent.), and U. S. P. (5.8 per cent.).

The mixture is known to be boiled long enough when a little of the clear liquid, poured into another test-tube and warmed, gives no effervescence on the addition of an acid (sulphuric, hydrochloric, or acetic)—a test whose mode of action will be explained hereafter.

*Best Method of expressing Decompositions.*—This will be easy of comprehension if what has already been stated concerning symbols and formulæ, on pages 31 to 41, has been carefully and thoughtfully considered. The best means of showing on paper the action which occurs when chemical substances attack each other is by the employment either of equations or diagrams, setting forth the formulæ of the molecules concerned in the reaction. In an *equation* the formulæ of the salts used are written on one line, the sign of addition (+) intervening; the sign of equality (=) follows, and then the formulæ of the salts produced, also separated by a plus sign (+). Thus:—

$$K_2CO_3 + Ca2HO = 2KHO + CaCO_3.$$

In this reaction (the operation just performed) the metals of the molecules of the two salts change place: from $K_2CO_3$ and Ca2HO there are produced $CaCO_3$ and KHO (two molecules, 2KHO); from carbonate of potassium and hydrate of calcium there result carbonate of calcium (the insoluble portion) and hydrate of potassium (in solution).*

* If the student is already accustomed to the use of ordinary equations, he may pass on to *Note* 1, on page 65. If not, the author would strongly recommend the temporary employment of diagrams for expressing chemical changes. Indeed, the occasional, if not the regular, use of graphic equations or diagrams is of advantage to all students. For while equation or diagram equally well records the formulæ of the salts concerned in the whole reaction, the diagram alone suggests the *mode* in which its writer believes the respective atoms to change

In constructing a *diagram* or pictorial illustration of a chemical reaction (the reaction, for instance, just described), *first* the formulæ of the salts used are written under each other on the left side of the leaf of a note-book, thus:—

$K_2CO_3$

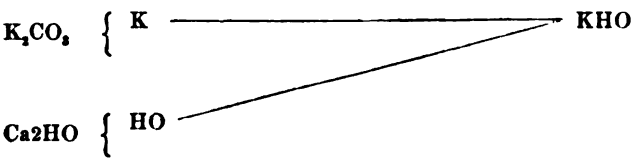
Ca2HO

Such formulæ are, in this Manual, always given with the description of the reaction. *Secondly*, on the right is then written the formula of the chief substance produced, thus:—

$K_2CO_3$ KHO

Ca2HO

*Thirdly*, the formation of this chief body under consideration, that is to say, both the origin of its elements and their destination, is traced out by the help of brackets which show the source of the elements, and converging lines which suggest the approach and final union of the elements, thus:—

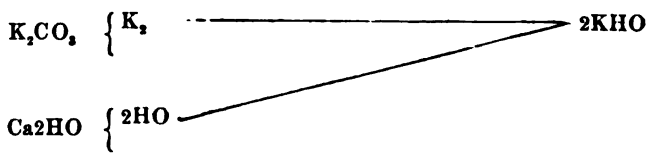



At this stage (at other stages, perhaps, in other reactions) the reader's own intelligent power of thought and reflection must come into exercise. He must reason somewhat as follows: "I am converting, and entirely converting, a quantity of carbonate of potassium into hydrate of potassium. A molecule, the smallest quantity I can picture on paper, of the carbonate of potassium ($K_2CO_3$) contains, I am told, two atoms of potassium ($K_2$), and a molecule of the hydrate (KHO), one atom (K). Therefore each molecule of the carbonate ($K_2CO_3$) will furnish two molecules of the hydrate (2KHO). Moreover, I notice that in the formula of a molecule of the hydrate of calcium (slaked lime) I employ, there are 2 of the HO (that is, 2HO), and this fact confirms me in the deduction that one molecule

---

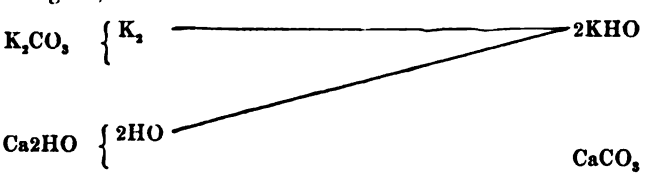
their positions. In the paragraphs succeeding the above, detailed explanations are given respecting the use and construction of diagrams.

of the carbonate affords two molecules of the hydrate." The pupil will then amend his diagram, thus:—

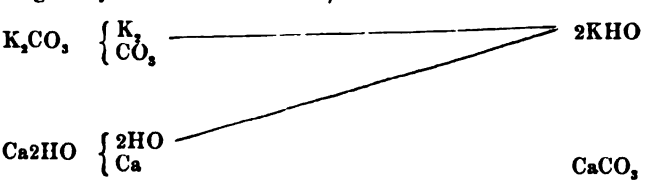



*Fourthly*, the question as to what becomes of the other elements must be cleared up. Indeed, when the reader remembers that he is studying this reaction for the aid it affords him in learning chemistry, and not because he is desirous of manufacturing caustic potash, he will see that this latter part of the reaction is quite as important as the former. To complete the diagram, then, he must first know what other compound is produced and its formula. The context of his Manual will generally afford this information, or, after a little experience is acquired, analogies or his own knowledge will suggest correct formulæ. In this case carbonate of calcium is produced, $CaCO_3$. (This product is, in fact, precipitated chalk, together with any excess of slaked lime and any natural impurities in the slaked lime. *Pure* "precipitated chalk" is made by an analogous reaction described subsequently.)

The source of the elements of the carbonate of calcium and, finally, their union, must be indicated just as the source and mode of formation of the potash were indicated. That is to say, after the formula of this second substance produced ($CaCO_3$) is written on the right of the diagram, thus:—

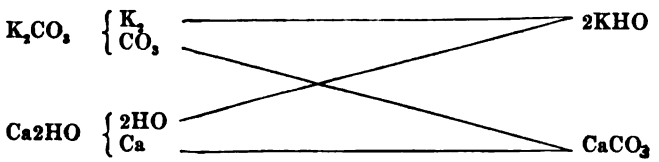



the source of its elements is shown by writing the symbols for those elements on the right of the bracket attached to the formula containing the symbols of the elements, thus:—




Lines converging from the symbols of the elements and uniting at the formula of the substance are then drawn, to suggest approach of

the atoms of the elements and their union to form a molecule of the compound. The diagram will now be complete, thus:—




The formation of a third product or a fourth product would be indicated in a similar manner.

*Note* 1.—It will be seen that the chief data required in making either equationary or diagrammatic notes of decompositions are the symbolic formulæ of the various compounds employed and produced. These formulæ are, in this Manual, given whenever necessary. Chemists obtain them in the first instance by help of quantitative analysis. By the same means is obtained a check on the probabilities respecting the relative number of molecules concerned in a reaction.

*Note* 2.—While an equation or a diagram is an attempt to picture the reaction which ensues when molecules of different substances act upon one another, it necessarily only represents two or a minimum number of the molecules. The student will, of course, understand that what is true of these two or three molecules is true of the thousands or millions of molecules forming the mass or whole quantity of material on which he experiments.

*Note on Nomenclature.—Hydrates* are bodies indirectly or directly derived from water by one-half of its hydrogen becoming displaced by an equivalent quantity of another radical. Thus, a piece of potassium thrown on to water (HHO) instantly liberates hydrogen, hydrate of potassium (KHO) being formed. The temperature produced at the same time is sufficiently high to cause ignition of the hydrogen, which burns with a purple flame (owing to the presence of a little vapor of potassium), while the hydrate of potassium remains dissolved in the bulk of the water. This radical or root or group of elements (HO), common to all hydrates, is sometimes termed *hydroxyl.* Water might be termed hydrate of hydrogen or hydroxylide of hydrogen.

*Explanation.*—With regard to the group of atoms represented by the symbols $CO_3$ and HO, only a few words need be said here. The former ($CO_3$) is the grouping (root or radical) found in all carbonates; it is termed the carbonic radical, and is as characteristic of carbonates as potassium (K) is of potassium salts. HO (hydroxyl) is characteristic of all hydrates. $CO_3$ is a bivalent root, HO is univalent; hence $CO_3$ is found

united with two univalent atoms, as in carbonate of potassium, $K_2CO_3$, or with one bivalent atom, as in carbonate of calcium, $CaCO_3$; and HO is found united in single proportion with univalent atoms, as in hydrate of potassium, KHO, or in double proportion with bivalent atoms, as in hydrate of calcium, Ca2HO. The quantivalence of a metal has only to be learnt, and the formulæ of its carbonate and hydrate are ascertained without seeing the formula of either. The formulæ of all other metallic salts are constructed on the same principle. But, beyond committing to memory the formulæ and qantivalence of the various groupings characteristic of carbonates, hydrates, nitrates, sulphates, acetates, etc. (see the following Table), special attention should not at present be devoted to the subject of the constitution of salts, but restricted to what may be called the metallic or basylous side of salts. The formulæ and quantivalence of the chief acidulous groupings referred to and the symbols and quantivalence of allied elementary bodies are included in the following Table:—

*Formulæ and Quantivalence of Acidulous Radicals.*

| | | |
|---|---|---|
| All chlorides contain . . . . | Cl | Univalent radicals. |
| " bromides " . . . . | Br | |
| " iodides " . . . . | I | |
| " cyanides " . . . . | CN | |
| " hydrates " . . . . | HO | |
| " nitrates " . . . . | $NO_3$ | |
| " chlorates " . . . . | $ClO_3$ | |
| " acetates " . . . . | $C_2H_3O_2$ | |
| " oxides " . . . . | O | Bivalent radicals. |
| " sulphides " . . . . | S | |
| " sulphites " . . . . | $SO_3$ | |
| " sulphates " . . . . | $SO_4$ | |
| " carbonates " . . . . | $CO_3$ | |
| " oxalates " . . . . | $C_2O_4$ | |
| " tartrates " . . . . | $C_4H_4O_6$ | Trivalent radicals. |
| " citrates " . . . . | $C_6H_5O_7$ | |
| " phosphates " . . . . | $PO_4$ | |
| " borates " . . . . | $BO_3$ | |

*Radicals.*—The above elements and compounds are termed *radicals*, each being the common *root* (*radix*) in a series of salts. Why compound radicals (as $NO_3$, $SO_4$, $PO_4$, etc.) differ in quantivalence cannot well be explained. Their constituent atoms doubtless always exert the same amount of attractive force, nearly but not quite all this force being exerted in re-

taining the atoms in one group, and the remainder probably determining the quantivalence. Some of the compound radicals are obtainable in the free state, others have yet to be proved capable of isolated existence.*

*Pure Solution of Potash.*—Solution of potash generally contains a trace of alumina dissolved from the lime by the hot alkali, but not enough to interfere with the use of the liquid in medicine. If the solution is required for analytical purposes, it may be obtained free from alumina by avoiding the employment of heat, in the manner suggested by Redwood. Half a gallon is made by mixing half a pound of slaked lime with about three pints of water, placing the mixture in a half-gallon bottle (Winchester quart), and adding to it, in small quantities at a time, a solution of half a pound of carbonate of potassium dissolved in the other pint of water, shaking the mixture well for several minutes after each addition. The whole is now set on one side till clear; and then, if a small quantity poured into a test-tube and warmed does not effervesce on the addition of hydrochloric acid, the solution is fit for use. If effervescence (due to carbonic acid gas) occurs, the mixture must be again well shaken. If the lime be good and recently slaked, and the bottle violently shaken once every half-hour, the decomposition will be complete in about ten or twelve hours.

*Liquor Potassæ* is officially directed to be made as follows:—

Dissolve 1 pound of carbonate of potassium in one gallon of water; heat the solution to the boiling-point in a clean iron vessel, gradually mix with it 12 ounces of slaked lime, and continue the ebullition for ten minutes with constant stirring. Then remove the vessel from the fire; and when by the subsidence of the insoluble matter the supernatant liquor has become perfectly clear, transfer it, by means of a siphon or by decantation, to a green-glass bottle, furnished with an air-tight stopper, and add distilled water, if necessary, to make it correspond with the tests of *specific gravity* and *neutralizing power*. (These tests and the mode of applying them will be explained in subsequent sections.)

*Solid Potash.*—Solution of potash evaporated to dryness in a silver or clean iron vessel and the residue fused and poured into moulds constitutes *Potassa Caustica*, B. P.; *Potassa*, U. S. P. It often contains chlorides, detected by nitrate of silver, and sulphates, detected by a barium salt, as described subsequently in connection with hydrochloric and sulphuric acids. *Potassa cum Calce*, U. S. P., is a grayish-white powder, made by rubbing together equal weights of solid potash and quicklime.

---

* Some modern authors term these roots *radicles*, a word more usefully expressive of little roots or rootlets.

## Sulphurated Potash.

*Second Synthetical Reaction.*—Into a test-tube put a few grains of carbonate of potassium previously mixed with half its weight of sulphur. Heat the mixture gradually until it ceases to effervesce. The resulting fused mass poured on a slab and quickly bottled is the *Potassa Sulphurata*, Sulphurated Potash, of B. P., or the *Potassii Sulphuretum*, U. S. P.

$$\underset{\text{Carbonate of potassium.}}{3K_2CO_3} + \underset{\text{Sulphur.}}{4S_2} = \underset{\text{Hyposulphite of potassium.}}{K_2S_2O_3} + \underset{\text{Sulphide of potassium.}}{2K_2S_3} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

As met with in pharmacy this salt is not a single definite chemical compound, but a mixture of several; in short, its chemical character is well indicated by its vague name. When fresh, and if carefully prepared with the official proportions of dry ingredients, it is of the color of liver (whence the old name "liver of sulphur"), and consists, as shown by J. Watts, of the salts mentioned in the foregoing equation, together with a little undecomposed carbonate of potassium, with perhaps higher sulphides of potassium ($K_2S_4$ and $K_2S_5$); but rapidly absorbing oxygen from the air it soon becomes green and yellow, sulphite ($K_2SO_3$) and sulphate of potassium ($K_2SO_4$) are formed, and ultimately a useless mass of dirty-white color results, consisting of sulphate and hyposulphite, with generally some carbonate of potassium and free sulphur. Moreover, if overheated in manufacture, the hyposulphite $4(K_2S_2O_3)$ is decomposed into sulphate $3(K_2SO_4)$ and sulphide ($K_2S_3$) of potassium. Recently made, "about three-fourths of its weight are dissolved by rectified spirit." It is occasionally employed in the form of ointment.

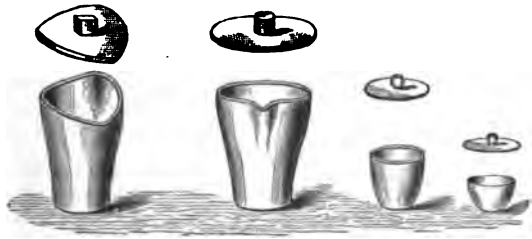
The extremely, indeed most unusually, complicated nature of the reaction, will probably cause failure to any attempt by a student to draw out an equation or a diagram of the reaction without the aid of the printed equation given above. He may therefore content himself, in this case, by introducing into his note-book a diagram founded directly on the equation and on the numbers of molecules there stated. With this printed equation, and the details of construction of diagrams given in connection with the first synthetical reaction, he will be able to give a diagram of this second reaction without troubling his reasoning powers, while at the same time he will be familiarizing himself with the more mechanical portions of a diagram.

In preparing large quantities of sulphurated potash, the test-tube is replaced by an earthenware vessel termed a *crucible* (from *crux*, a cross, for originally a cross was impressed upon the melting-pot as used by alchemists and goldsmiths; others derive the word from crux, an instrument of torture, the sense here being symbolical).

*Heating Crucibles.*—Crucibles of a few ounces' capacity may be heated in an ordinary grate-fire. Larger ones require a stove with a good draught—that is, a *furnace*. Even the smaller ones are more conveniently and quickly heated in a furnace. Half-ounce or one

ounce experimental porcelain crucibles may be heated in a spirit- or gas-flame; the air-gas flame already described being generally the most suitable.

Fig. 12. Fig. 13. Fig. 14.



Crucibles of various forms.

## Acetate of Potassium.

*Third Synthetical Reaction.*—Place ten or twenty or more grains of carbonate of potassium in a small dish, and *saturate* (*satur*, full) with acetic acid; that is, add acetic acid so long as effervescence is thereby produced; the resulting liquid is a strong solution of acetate of potassium.

Evaporate most of the water in an open dish (See Figs. 15, 16), stirring with a glass rod* to promote the evolution of vapor; a white salt remains, which fuses on the further application of heat; this is the official Acetate of Potassium (*Potassæ Acetas*, B. P., and *Potassii Acetas*, U. S. P.). If fused in the open vessel the acetate is liable to become slightly charred and discolored; this is prevented by transferring the solid residue to a test-tube or Florence flask before finally fusing. It forms a white, deliquescent, foliaceous, satin-like mass, neutral to test-paper, and wholly soluble in spirit. A ten per cent. solution in water forms the "Solution of Acetate of Potash," B. P.

$$\underset{\text{Carbonate of potassium.}}{K_2CO_3} + \underset{\text{Acetic acid.}}{2HC_2H_3O_2} = \underset{\text{Acetate of potassium.}}{2KC_2H_3O_2} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

*Explanation of Formulæ.*—The formula for one molecule of acetic acid (the acetate of hydrogen) is $HC_2H_3O_2$, and one of acetate of potassium, $KC_2H_3O_2$. The grouping, $C_2H_3O_2$, is characteristic of

* Glass rod is usually purchased in the form of long sticks. The pieces may be cut to convenient lengths of from 6 to 12 inches (*vide* p. 16), sharp ends being rounded off by holding in a flame for a few minutes.

all acetates; it is univalent, and may be shortly, though less instructively written $\overline{A}$.

*Explanation of Process.*—When two molecules of acetic acid ($2HC_2H_3O_2$) and one of carbonate of potassium ($K_2CO_3$) react, two molecules of acetate of potassium ($2KC_2H_3O_2$) and one of carbonic acid ($H_2CO_3$) are produced, the latter at once splitting up into water ($H_2O$) and carbonic acid gas ($CO_2$), as already shown in the equation.

*Diagram of the Reaction.*—The nature of the above operation is indicated by an *equation;* it (and succeeding reactions) should be expressed in the student's note-book as a *diagram*, and, if possible, without the aid of the equation.

*Note.*—The above reaction has a general as well as a special synthetical interest. It represents one of the commonest methods of forming salts, namely, the saturation of an acid with a carbonate. Carbonates added to acetic acid yield acetates, to nitric acid nitrates, to sulphuric acid sulphates. Many illustrations of this general process occur in pharmacy.

*Evaporation* of water from a liquid is best conducted in wide shallow vessels rather than in narrow deep ones, as the steam can thus quickly diffuse into the air and be rapidly conveyed away; hence a small round-bottomed basin, heated as shown in Fig. 15, is far more suitable than a test-tube for such operations. On the manufacturing scale, iron, or iron lined

Fig. 15. Fig. 16.

Evaporation from small and large basins.

with enamel, or semi-porcelain, copper, tinned copper, or solid tin pans are used. Up to 12 or 18 inches diameter, pans, basins, or dishes, made of Wedgwood ware or porcelain composition (Fig. 16), may be employed. Small dishes may be supported by retort-stands (Fig. 15), larger by cylinders (Fig. 16), to which the dish is, if less in diameter than the cylinder,

adapted by such flat rings or diaphragms as are shown in the figure.

## Bicarbonate of Potassium.

*Fourth Synthetical Reaction.*—Make a *strong* solution of carbonate of potassium by heating in a test-tube a mixture of several grains of the salt with rather less than an equal weight of water. Through the cooled solution pass carbonic acid gas slowly but continuously; after a time a white crystalline precipitate of Acid Carbonate or Bicarbonate of Potassium ($KHCO_3$), *Potassii Bicarbonas*, U. S. P., the Bicarbonate of Potash of the British Pharmacopœia (*Potassæ Bicarbonas*, B. P.), will be formed.

$$\underset{\text{Carbonate of potassium.}}{K_2CO_3} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2} = \underset{\text{Bicarbonate of potassium.}}{2KHCO_3}$$

*The carbonic acid* gas necessary for this operation is to be prepared from marble, though it might be obtained from any carbonate. Thus the previous synthetical reaction could be made available for this purpose, the carbonic acid gas evolved on the addition of the acetic acid to the carbonate of potassium being conducted into a strong solution of more carbonate of potassium by a glass tube bent and fitted as described when treating of oxygen gas. But motives of economy induce the use of carbonate of calcium, the form known as marble being always employed. Economy and convenience also cause hydrochloric acid to be used in preference to acetic or any other.

*Generate the carbonic acid gas* by adding common hydrochloric acid, diluted with twice its bulk of water, to a few fragments of marble contained in a test-tube or small flask, and conduct the gas into the solution of carbonate of potassium by a glass tube bent to a convenient angle or angles, and fitted to the test-tube by a cork in the usual way. (See Fig. 10, though no heat is necessary.) The tube may be replenished with marble, or acid, or both, when the evolution of gas is becoming slow. In working on any larger quantity than a few grains of the carbonate, a wide delivery tube should be employed, or the end of the narrow tube occasionally cleared from any bicarbonate that may have been deposited in it. The more economical official arrangements of the apparatus employed in this process will be described under the corresponding sodium salt.

*Deposition of the Bicarbonate explained.*—Bicarbonate of potassium is to a certain extent soluble in water; but as it is less so than the carbonate of potassium, and as a *saturated* solution of the latter has been used, the precipitation of a part of the bicarbonate inevita-

bly occurs. In other words, the quantity of water present is sufficient to keep the carbonate, but insufficient to retain the equivalent quantity of bicarbonate in solution.

*Properties.*—Prepared on the large scale, bicarbonate of potassium occurs in colorless, non-deliquescent, right rhombic prisms; it has a saline, feebly alkaline, non-corrosive taste.

*Effervescing Solution of Potash.*—A solution of 30 grains of bicarbonate of potassium in one pint of water, charged with 7 times its bulk (often less) of carbonic acid gas by pressure, constitutes the ordinary "potash-water," the so-called *Liquor Potassæ Effervescens*, B. P.

*Notes on Nomenclature.*—The prefix *bi-* in the name "bicarbonate of potassium" serves to recall the fact that to a given amount of potassium this salt contains *twice* as much carbonic radical as the carbonate. The salt is really a "carbonate of potassium and hydrogen" ($KHCO_3$); it is intermediate between carbonate of potassium ($K_2CO_3$) and carbonate of hydrogen, or true carbonic acid ($H_2CO_3$); it is "acid carbonate of potassium" or "hydric potassium carbonate;" in chemical constitution it is an acid salt, although not acid to the taste.

Salts whose specific names end in the syllable "*ate*" (carbon*ate*, sulph*ate*, etc.) are in general conventionally so termed when they contain an acidulous radical, or the characteristic elements of an acid, whose name ends in "*ic*," and from which acid they have been or may be formed. Thus the syllable "*ate*," in the words sulph*ate*, nitr*ate*, acet*ate*, carbon*ate*, etc., indicates that the respective salts contain a radical whose name ended in *ic*, the previous syllables, sulph-, nitr-, acet-, carbon-, indicating what that radical was—the sulphuric, nitric, acetic, or carbonic. Occasionally a letter or syllable is dropped from or added to a word to render the name more euphonious; thus the sulphuric radical forms sulphates, not sulphurates.

## Citrate of Potassium.

*Fifth Synthetical Reaction.*—Dissolve a few grains or more of carbonate (bicarbonate, U. S. P.) of potassium in water, and add citric acid ($H_3C_6H_5O_7$) until it no longer causes effervescence. The resulting liquid is a solution of citrate of potassium ($K_3C_6H_5O_7$) (*Liquor Potassii Citratis*, U. S. P.). Evaporated to dryness, in an open dish, a pulverulent or granular residue is obtained, which is the official *Potassii Citras*, U. S. P. (*Potassæ Citras*, B. P.), a white deliquescent powder.

$$\underset{\text{Carbonate of potassium.}}{3K_2CO_3} + \underset{\text{Citric acid.}}{2H_3C_6H_5O_7} = \underset{\text{Citrate of potassium.}}{2K_3C_6H_5O_7} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

*Citrates.*—The citric radical or group of elements, which with three atoms of hydrogen forms a molecule of citric acid, and with three of potassium citrate of potassium, is a trivalent grouping; hence the three atoms of potassium in a molecule of the citrate. The

full chemistry of citric acid and other citrates will be subsequently described.

*Nitrate of potassium* ($KNO_3$) (*Potassæ Nitras*, B. P.; *Potassii Nitras*, U. S. P.) and *Sulphate of potassium* ($K_2SO_4$) (*Potassæ Sulphas*, B. P.; *Potassii Sulphas*, U. S. P.) could obviously also be made by saturating nitric acid ($HNO_3$), and sulphuric acid ($H_2SO_4$), respectively, by carbonate of potassium. Practically they are not made in that way—the nitrate occurring, as already stated, in nature, and the sulphate as a by-product in many operations. Both salts will be hereafter alluded to in connection with nitric acid.

## Tartrate of Potassium.

*Sixth Synthetical Reaction.*—Place a few grains of carbonate of potassium in a test-tube with a little water, heat to the boiling-point, and then add acid tartrate of potassium ($KHC_4H_4O_6$ or $KH\overline{T}$) till there is no more effervescence; a solution of neutral tartrate of potassium ($K_2\overline{T}$) results, the *Potassii Tartras* of the United States Pharmacopœia (*Potassæ Tartras*, B. P.), the old "Soluble Tartar." Crystals (4- or 6-sided prisms) may be obtained on concentrating the solution by evaporation and setting the hot liquid aside. Larger quantities are made in the same way, 20 of acid tartrate and 9 of carbonate (with 50 of water) being about the proportions necessary for neutrality.

$$\underset{\text{Acid tartrate of potassium.}}{2KHC_4H_4O_6} + \underset{\text{Carbonate of potassium.}}{K_2CO_3} = \underset{\text{Neutral tartrate of potassium.}}{2K_2C_4H_4O_6} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

*Tartrates.*—$C_4H_4O_6$ are the elements characteristic of all tartrates; they form a bivalent grouping; hence the formula of the hydrogen tartrate, or tartaric acid, is $H_2C_4H_4O_6$; that of the potassium tartrate $K_2C_4H_4O_6$; of the intermediate salt, the acid potassium tartrate (cream of tartar), $KHC_4H_4O_6$. If the acid tartrate of one metal and the carbonate of another react, a neutral dimetallic tartrate results, as seen in Rochelle salt ($KNaC_4H_4O_6$), the *Soda Tartarata* of the British Pharmacopœia. (*Potassii et Sodii Tartras*, U. S. P.)

*Acid salts* (*e. g.* $KHC_4H_4O_6$), that is, salts intermediate in composition between a normal or neutral salt (*e. g.* $K_2C_4H_4O_6$) and an acid (*e. g.* $H_2C_4H_4O_6$) will frequently be met with. All acidulous radicals, except those which are univalent, may be concerned in the formation of acid salts.

## Iodide of Potassium.

*Seventh Synthetical Reaction.*—To a solution of potash, heated in a test-tube, flask, or evaporating-basin, according to quantity, add a small quantity of solid iodine. The deep color

of the iodine disappears entirely. This is due to the formation of the colorless salts, iodide of potassium (KI) and iodate of potassium ($KIO_3$), which remain dissolved in the liquid. Continue the addition of iodine so long as its color, after a few minutes' warming and stirring, disappears. When the whole of the potash in the solution of potash has been converted into the salts mentioned, the slight excess of iodine remaining in the liquid will color it, and thus show that this stage of the operation is completed.

$$\underset{\text{Hydrate of potassium.}}{6KHO} + \underset{\text{Iodine.}}{3I_2} = \underset{\text{Iodide of potassium.}}{5KI} + \underset{\text{Iodate of potassium.}}{KIO_3} + \underset{\text{Water.}}{3H_2O}$$

*Separation of the Iodide from the Iodate.*—Evaporate the above solution to dryness. If each salt were required, the solid mixture might be digested in spirit of wine, which dissolves the iodide, but not the iodate. But the iodide only is used in medicine. Mix the residue, therefore (reserving a grain or two for a subsequent experiment), with about a twelfth of its weight of charcoal, and gently heat in a test-tube or crucible until slight deflagration ensues.*

The crucible may be held in a spirit or air-gas flame, or other fire, by tongs. (Scissors-shaped and other "crucible-tongs" are sold by all makers of apparatus.) Under these circumstances the iodide remains unaffected; but the iodate loses all its oxygen, and is thus also reduced to the state of iodide.

$$\underset{\text{Iodate of potassium.}}{2KIO_3} + \underset{\text{Carbon.}}{3C_2} = \underset{\text{Iodide of potassium.}}{2KI} + \underset{\text{Carbonic oxide.}}{6CO}$$

Treat the mass with a little water, and filter to separate excess of charcoal; a solution of pure iodide of potassium results. It may be used as a reagent or evaporated to a small bulk and set aside to crystallize.

This is the process mentioned in the British and United States Pharmacopœias (*Potassii Iodidum.*) "Solution of Iodate of Potassium" is also official as a test-liquid.

---

* If, in the operation of heating iodate of potassium with charcoal, excess of the latter be employed, slight incandescence rather than deflagration occurs; if the charcoal be largely in excess, the reduction of the iodate to iodide of potassium is effected without visible deflagration or even incandescence.

Deflagration means violent burning, from *flagratus*, burnt (*flagro*, I burn), and *de*, a prefix augmenting the sense of the word to which it may be attached. Paper thrown into a fire simply burns, nitre deflagrates. *De*-tonate (*detono*) is a precisely similar word, meaning to explode with violent noise.

*Properties.*—Iodide of potassium crystallizes in small cubical crystals, very soluble in water, less so in spirit. One part in ten of water forms "Solution of Iodide of Potassium," B. P. Exposed to air and sunlight, pure iodide of potassium becomes slightly brown owing to the liberation of iodine. A little carbonate of potassium is produced by action of the atmospheric carbonic acid and a little hydriodic acid (HI) is set free: the latter, attacked by oxygen, yielding a trace of water and of free iodine. The ozone in the air (see "Ozone" in Index) may also contribute to the liberation of iodine from such compounds as iodide of potassium.

*The addition of charcoal* in the above process is simply to facilitate the removal of the oxygen of the iodate of potassium. Iodate of potassium ($KIO_3$) is analogous in constitution, and in composition, so far as the atoms of oxygen are concerned, to chlorate of potassium ($KClO_3$), which has already been stated to be more useful than any other salt for the actual preparation of oxygen gas itself. Hence the removal of the oxygen of the iodate might be accomplished by heating the residue without charcoal. In that case the liberated oxygen would be detected on inserting the incandescent extremity of a strip of wood into the mouth of the test-tube in which the mixture of iodide and iodate had been heated. The charcoal, however, burns out the oxygen more quickly, and thus economizes heat and time.

*Note.*—The formula of iodide of potassium (KI) shows that the salt contains potassium and iodine in atomic proportions. A reference to the table of atomic weights at the end of the volume, and a rule-of-three sum, would therefore show what weight of salt is producible from any given weight of iodine.

*Detection of Iodate in Iodide of Potassium.*—Iodate of potassium remaining as an impurity in iodide of potassium may be detected by adding to a solution of the latter salt some weak acid (say tartaric), shaking, and then adding mucilage of starch; blue "iodide of starch" is formed if a trace of iodate be present, not otherwise. By the reaction of the added acid and the iodate of potassium, iodic acid ($HIO_3$) is produced, and by reaction of the added acid and the iodide of potassium, hydriodic acid (HI) is produced; neither of these two acids alone attacks starch, but by reaction on each other the two give rise to free iodine, which then forms the blue color. This experiment should be tried on a sample of pure iodide of potassium and on a grain or two of the impure iodide reserved from the previous experiment.

$$HIO_3 + 5HI = 3H_2O + 3I_2$$

*Note on Nomenclature.*—The syllable *ide* attached to the syllable *iod*, in the name "iodide of potassium," indicates that the element *iodine* is combined with the potassium. An iod*ate*, as already explained, is a salt containing the characteristic elements of iod*ic* acid and all iodic compounds. Salts one of whose names ends in *ide* are those which are, or may be, formed from elements. The names of

salts which are, or may be, formed from compounds include other syllables, *ate* being one (see page 72). The only other syllable is *ite*, which is included in the names of salts which are, or may be, formed from acids and radicals whose names end in *ous:* thus hypo-sulph*ite* of sodium, etc. To recapitulate: A salt, whose name ends in *ate* contains a compound acidulous radical whose name ends in *ic;* a salt whose name ends in *ite* contains a compound acidulous radical whose name ends in *ous;* a salt whose name ends in *ide* contains an element for its acidulous radical. Thus sulph*ide* relates to sulphur, sulph*ite* to the sulphurous radical, sulph*ate* to the sulphuric radical, and so on with other "ides," "ites," or "ates."

*Bromide of Potassium* (*Potassii Bromidum*, B. P.).—This salt is identical in constitution with iodide of potassium, and may be made in exactly the same way, bromine being substituted for iodine. The formula of bromic acid is $HBrO_3$. It will be noticed that the following equations are similar in character to those showing the preparation of iodide of potassium.

$$\underset{\text{Hydrate of potassium.}}{6KHO} + \underset{\text{Bromine.}}{3Br_2} = \underset{\text{Bromide of potassium.}}{5KBr} + \underset{\text{Bromate of potassium.}}{KBrO_3} + \underset{\text{Water.}}{3H_2O}$$

$$\underset{\text{Bromate of potassium.}}{2KBrO_3} + \underset{\text{Carbon.}}{3C_2} = \underset{\text{Bromide of potassium.}}{2KBr} + \underset{\text{Carbonic oxide.}}{6CO}$$

*Potassii Bromidum*, U. S. P., is made by decomposing solution of bromide of iron ($FeBr_2$) by solution of pure carbonate of potassium ($K_2CO_3$), evaporating, and crystallizing.

## Manganates of Potassium.

*Eighth Synthetical Reaction.*—Place a fragment of solid caustic potash (KHO), with about the same quantity of chlorate of potassium ($KClO_3$), and of black oxide of manganese ($MnO_2$) on a piece of platinum foil.* Hold the foil by a small pair of forceps or tongs in the flame of a blowpipe for a few minutes until the fused mixture has become dark green. This color is that of *manganate of potassium* ($K_2MnO_4$).

$$\underset{\text{Hydrate of potassium.}}{6KHO} + \underset{\text{Chlorate of potassium.}}{KClO_3} + \underset{\text{Black oxide of manganese.}}{3MnO_2} = \underset{\text{Manganate of potassium.}}{3K_2MnO_4} + \underset{\text{Chloride of potassium.}}{KCl} + \underset{\text{Water.}}{3H_2O}$$

*Ninth Synthetical Reaction.—Permanganate of Potassium.* ($K_2Mn_2O_8$) (*Potassæ Permanganas*, B. P., *Potassii Perman-*

* The foil may be 1 inch broad by 2 inches long. No ordinary flame will melt the platinum; fused caustic alkalies only slowly corrode it, and very few other chemical substances affect it at all; hence the same piece may be used in experiments over and over again. Most metals form a fusible alloy with platinum, and phosphorus rapidly attacks it, hence such substances, as well as mixtures likely to yield them, should be heated in a small porcelain crucible.

*ganas*, U. S. P.), which is purple, is obtained, or rather a solution of it, on placing the foil and its adherent mass in water, and boiling for a short time.

$$\underset{\text{Manganate of potassium.}}{3K_2MnO_4} + \underset{\text{Water.}}{2H_2O} = \underset{\text{Permanganate of potassium.}}{K_2Mn_2O_8} + \underset{\text{Hydrate of potassium.}}{4KHO} + \underset{\text{Black oxide of manganese.}}{MnO_2}$$

On the large scale, the potash set free in the reaction is neutralized by sulphuric or carbonic acid, and the solution evaporated to the crystallizing point. Further details will be given in connection with manganese.

Solutions of manganate or permanganate of potassium and of sodium so readily yield their oxygen to organic matter, that they are used on the large scale as disinfectants, under the name of "Condy's Disinfecting Fluids."

*Synthetical Reactions* bringing under consideration the remaining official compounds (namely, bichromate, arsenite, chlorate, cyanide, ferrocyanide, and ferridcyanide of potassium) are deferred at present.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*Note.*—These are reactions utilized in searching for small quantities of a substance (in the present instance of potassium) in a solution. They are best performed in test-tubes or other small vessels. Each should be expressed, in the form of an equation or diagram, in the student's note-book. *All previous or future equations given in this volume should be transferred to the note-book in the form of diagrams, constructed as described on pages* 63 *and* 64, *unless the student can with ease construct the equations without the aid of the manual.*

*First Analytical Reaction.**—To a solution of any salt of

* As already indicated, chemical reactions are scarcely analytical or synthetical in themselves, but, rather, performed with an analytical or synthetical object. Indeed not unfrequently one and the same reaction is both a synthetical and an analytical reaction. Thus this first, so-called, "analytical reaction" is a synthetical reaction if performed with the object of preparing a specimen of the double chloride of platinum and potassium. It is an analytical reaction, or, rather, has analytical interest, if performed with the object of demonstrating the presence of potassium. Chemical reactions in themselves are operations, not so much of analysis (resolution) or synthesis (combination) or of analysis and synthesis conjoined, as of what has sometimes been termed *metathesis* (transposition). Molecules are not torn to atoms in an operation performed with an analytical object, nor are the atoms put together or set together in an operation (perhaps the same operation) performed with a synthetical object: but in both operations the atoms of the molecules undergo metathesis, that is, exchange places, or are transposed. In short, chemists use the words "analytical" and "synthetical" in a conventional rather than a strictly etymological sense.

potassium (chloride,* for example) add a few drops of hydrochloric acid and of a solution of perchloride of platinum ($PtCl_4$), and stir the mixture with a glass rod; a yellow granular or slightly crystalline precipitate† slowly forms. (This precipitate is the double chloride of platinum and potassium, and its composition is expressed by the formula $PtCl_4 2KCl$.)

*Memoranda.*—Experiments with such expensive reagents as perchloride of platinum are economically performed with watch-glasses, drops of the liquid being operated on. When the precipitate is long in forming, it is sometimes of an orange-yellow tint. If iodide of potassium happen to be the potassium salt under examination, some iodide of platinum ($PtI_4$) will also be formed, giving a red color to the solution, and a larger quantity of the *precipitant* (that is, the precipitating agent) be required.

*Precaution.*—Only chloride of potassium forms this characteristic compound; hence, if the potassium salt in the solution is known not to be a chloride, or if its composition is unknown, a few drops of hydrochloric acid must be added, otherwise some of the perchloride of platinum will be utilized for its chlorine only, the platinum being wasted. Thus, if nitrate of potassium ($KNO_3$) be present, a few drops of hydrochloric acid enable the potassium to assume the form of chloride when the perchloride of platinum is added, nitric acid ($HNO_3$) being set free.

*Explanation.*—The precipitate is, practically, insoluble in water. It is for this reason that a very small quantity of any soluble potassium salt (or, rather, of the potassium in that salt) is thrown out of solution by perchloride of platinum.

*Note on Nomenclature.*—When distinct molecules of salts unite and form a single crystalline compound, the product is termed a *double* salt. The double chloride of potassium and platinum is such a body.

## Acid Tartrate of Potassium.

*Second Analytical Reaction.*—To a solution of any salt of potassium add excess of strong solution of tartaric acid ($H_2C_4H_4O_6$), and shake or well stir the mixture; a white granular precipitate of acid tartrate of potassium ($KHC_4H_4O_6$) will be formed.

*Note.*—By "excess" of any test liquid (such as the "solution of tartaric acid" just mentioned) is meant such a quantity as is *probably*

---

* A few fragments of carbonate of potassium, two or three drops of hydrochloric acid, and a small quantity of water, give a solution of chloride of potassium at once, $K_2CO_3 + 2HCl = 2KCl + H_2O + CO_2$.

† By *precipitation* (from *præcipito*, to throw down suddenly) is simply meant the formation of particles of solid in a liquid, no matter whether the solid, the *precipitate*, subsides or floats.

rather more than sufficient to convert the whole weight of the compound attacked into the compound produced. Thus, in the present case enough acid must be added to convert the whole of the potassium salt operated on into acid tartrate of potassium. What the weight of salt operated on was must be mentally estimated, roughly, by the operator. It is not necessary in analyzing to know the exact weights of salts employed. The analyst must use his judgment, founded on his knowledge of the reaction (as shown by an equation) and of the molecular weights of the substances employed in the reaction as well as by the rough estimate of the amount of material on which he is experimenting.

*Limits of the Test.*—Acid tartrate of potassium is soluble in about 180 parts of cold and in 6 parts of boiling water. Hence, in applying the tartaric test for potassium, the solutions must not be hot. Even if cold, no precipitate will be obtained if the solutions are very dilute. This test, therefore, is of far less value than the first mentioned. The acid tartrate of potassium is less soluble in diluted alcohol than in water; so that the addition of spirit of wine renders the reaction somewhat more delicate.

*Cream of Tartar.*—The precipitate is the *Bitartrate or Acid Tartrate of Potassium* (*Potassæ Tartras Acida*, B. P., *Potassii Bitartras*, U. S. P.), though the official preparation is not formed in the above manner; on the contrary, the acid is derived from the salt, which occurs naturally in the juice of many plants.

*Memorandum.*—When the tartaric acid is added to the salt of potassium, and the acid tartrate formed, the acid whose chief elements were previously with the potassium is set free; and in such acid solutions the acid tartrate is somewhat soluble. To prevent loss on this account, acid tartrate of sodium, a salt tolerably soluble in water, may be used as a test instead of tartaric acid (Plunkett). The sodium, uniting with the acidulous radical, thus gives a neutral instead of an acid solution. But this advantage is of less importance from the fact that more water is introduced by the saturated solution of acid tartrate of sodium than by a saturated solution of tartaric acid.

*Third Analytical Reaction.*—The *flame-test.* Dip the looped end of a platinum wire into a solution containing a potassium salt, and introduce the loop into the lower part of a spirit-flame, the flame of a mixture of gas and air, a blowpipe flame, or other slightly colored flame. A light violet or lavender tint will be communicated to the flame, an effect highly characteristic of salts of potassium.

*Fourth Analytical Fact.*—Salts of potassium are not volatile. Place a fragment of carbonate, nitrate, or any other potassium salt, on a piece of platinum foil, and heat the latter in the flame of a lamp; the salt may fuse to a transparent liquid and flow freely over the foil, water also if present will escape as steam, and black carbon be set free if the salt happen to be a tartrate, citrate, etc.; but the potassium compound

itself will not be vaporized. This is a valuable negative property, as will be evident when the analytical reactions of ammonium come under notice.

## QUESTIONS AND EXERCISES.

70. Name the sources of Potassium.
71. Give the source, formula, and characters of Carbonate of Potassium.
72. Distinguish between synthetical and analytical reactions.
73. How is the official *Liquor Potassæ* prepared?
74. What is the systematic name of Caustic Potash?
75. State the chemical formula of Caustic Potash.
76. Construct an equation or diagram expressive of the reaction between Carbonate of Potassium and slaked lime.
77. Define a *hydrate*.
78. What group of atoms is characteristic of all carbonates?
79. Define the term *radical*.
80. How is "Sulphurated Potash" made, and of what salts is it a mixture?
81. What is the formula of the acetic radical—the radical of all acetates?
82. Draw a diagram showing the formation of Acetate of Potassium.
83. Give a general process for the conversion of carbonates into other salts.
84. What is the difference between Carbonate and Bicarbonate of Potassium? How is the latter prepared?
85. What is the relation between salts whose specific names end in the syllable "*ate*," and acids ending in "*ic*"?
86. Draw out diagrams or equations descriptive of the formation of Tartrate of Potassium from the Acid Tartrate, and Citrate from the Carbonate of Potassium.
87. Distinguish between a normal and an acid salt.
88. How is Iodide of Potassium made? Illustrate the process either by diagrams or equations.
89. Describe the appearance and chemical properties of Iodide of Potassium.
90. How much Iodide of Potassium is producible from 1000 grains of Iodine? Ans. 1307 grains.
91. Give a method for the detection of Iodate in Iodide of Potassium. Explain the reaction.
92. Has the syllable "*ide*" any general significance in chemical nomenclature?
93. What is the difference between sulphides, sulphites, and sulphates?
94. Mention the chemical relations of Bromide to Iodide of Potassium.
95. Describe the formation of Permanganate of Potassium, giving equations or diagrams.

96. How do manganate and permanganate of potassium act as disinfectants?

97. Enumerate the tests for potassium, explaining by diagrams or equations the various reactions which occur.

---

## SODIUM.

Symbol Na. Atomic weight 23.

Formula $Na_2$. Probable molecular weight 46.

*Memoranda.*—Most of the sodium salts met with in Pharmacy are obtained directly from carbonate of sodium which is now manufactured on an enormous scale from chloride of sodium (common salt, sea-salt or bay-salt, or rock-salt), the natural source of the sodium salts. When pure, salt (*Sodii Chloridum*, B. P. and U. S. P.) occurs "in small white crystalline grains, or transparent cubic crystals, free from moisture;" the best varieties commonly contain a little chloride of magnesium and sometimes other impurities. Besides the direct and indirect use of carbonate of sodium, or carbonate of soda, as it is commonly called in medicine, it is largely used for household cleansing purposes under the name of "soda," and in the manufacture of soap. Nitrate of sodium also occurs in nature, but is valuable for its nitric constituents rather than its sodium. Sodium is a constituent of about forty chemical or Galenical preparations of the Pharmacopœias.

Sodium is prepared by a process similar to that for potassium, but with less difficulty. Its atom is univalent, Na'.

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Reactions having Synthetical Interest.*

#### Hydrate of Sodium. Caustic Soda.

*First Synthetical Reaction.*—The formation of solution of hydrate of sodium or caustic soda, NaHO (*Liquor Sodæ*, B. P. and U. S. P.). This operation resembles that of making solution of potash already described.

The practical student should refer to the remarks made concerning solution of potash, applying them to solution of soda. He may perform the corresponding experiments or omit them, as he considers he does or does not clearly comprehend all they are designed to teach.

$$\underset{\text{Carbonate of sodium.}}{Na_2CO_3} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Hydrate of sodium.}}{2NaHO} + \underset{\text{Carbonate of calcium.}}{CaCO_3}$$

*Pure Solution of Soda*, free from any trace of alumina may be prepared by shaking in a Winchester quart, once every 20 or 30 minutes for 5 or 6 hours, 14 ozs. of crystals of carbonate of sodium and 8 ozs. of good recently slaked lime. The official *Liquor Sodæ* is made from 28 ounces of crystals of carbonate of sodium, 12 of slaked lime, and 1 gallon of water, under precisely similar circumstances to those detailed for *Liquor Potassæ* (p. 67). If the solution be evaporated to dryness, and the residue fused and poured into moulds, solid hydrate of sodium (*Soda Caustica*, B. P., *Soda*, U. S. P.) is obtained. Common and cheap caustic soda is now largely employed in manufactures. It is a by-product in the preparation of carbonate of sodium, and, though highly useful as a chemical agent, is too impure for use in medicine.

*Action of Sodium on Water.*—Sodium, like potassium, decomposes water with production of hydrate of sodium and hydrogen, but unless the sodium is confined to one spot, by placing it on a small floating piece of filter paper, the action is not sufficiently intense to cause ignition of the escaping hydrogen. When the latter does ignite, it burns with a yellow flame, due to the presence of a little vapor of sodium.

*Second Synthetical Reaction.*—The reaction of sulphur and carbonate of sodium at a high temperature resembles that of sulphur and carbonate of potassium; but, as the product is not used in medicine, the experiment may be omitted. It is mentioned here to draw attention to the close resemblance of the potassium salts to those of sodium.

## Acetate of Soda.

*Third Synthetical Reaction.*—Add the powder or fragments of carbonate of sodium ($Na_2CO_3$) to some strong acetic acid in a test-tube or evaporating-basin as long as effervescence occurs, and then evaporate some of the water.* When the solution is cold, crystals of acetate of sodium ($NaC_2H_3O_2,3H_2O$) (*Sodæ Acetas*, B. P., *Sodii Acetas*, U. S. P.) will be deposited. A ten per cent. solution in distilled water forms the "Solution of Acetate of Soda," B. P.

Acetate of sodium effloresces in dry air, and loses all its water of crystallization when gently heated. It supports a temperature of 270° or 280° F. without decomposition, but above 300° soon chars.

$$\underset{\text{Carbonate of sodium.}}{Na_2CO_3} + \underset{\text{Acetic acid.}}{2HC_2H_3O_2} = \underset{\text{Acetate of sodium.}}{2NaC_2H_3O_2} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

* The "water" alluded to occurs in the acid, which, though commonly termed "acetic acid, is really a solution of that acid in water.

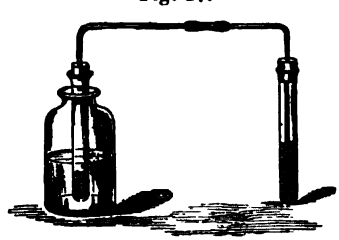
## Bicarbonate of Sodium.

*Fourth Synthetical Reaction.*—The action of carbonic acid ($H_2CO_3$), or carbonic acid gas ($CO_2$) and water ($H_2O$), on carbonate of sodium ($Na_2CO_3$). This resembles that of carbonic acid on carbonate of potassium, but is applied in a different manner. The result is bicarbonate of sodium ($NaHCO_3$) (*Sodæ Bicarbonas*, B. P., *Sodii Bicarbonas Venalis*, U. S. P.).

| $Na_2CO_3$ | + | $H_2O$ | + | $CO_2$ | = | $2NaHCO_3$ |
|---|---|---|---|---|---|---|
| Carbonate of sodium. | | Water. | | Carbonic acid gas. | | Bicarbonate of sodium. |

*Process.*—Heat crystals of carbonate of sodium in a porcelain crucible until no more steam escapes. Rub the product, in a mortar, with two-thirds its weight of more of the crystals and place the powder in a test-tube or small bottle into which carbonic acid gas may be conveyed by a tube passing through a cork and terminating at the bottom of the vessel. To generate the carbonic acid gas fill a test-tube having a small hole in the bottom (or a similar piece of glass tubing, of which one end is plugged by a grooved cork) with fragments of marble, insert a cork and delivery-tube, and connect the latter with the similar tube of the vessel containing the carbonate of sodium by a piece of India-rubber tubing. Now plunge the tube of marble into a test-glass, or other vessel, containing a mixture of one part hydrochloric acid and two parts water, and loosen the cork of the carbonate-of-sodium tube until carbonic acid gas, generated in the marble tube, may be considered to fill the whole arrangement; then replace the cork tightly and set the apparatus aside. As the gas is absorbed by the carbonate of sodium, hydrochloric acid rises into the marble tube, generates fresh gas, which, in its turn, drives back the acid liquid, and thus prevents the production of any more gas until further absorption has occurred. When the salt is wholly converted into bicarbonate ($NaHCO_3$), it will be found to have become damp through the liberation of water from the crystallized carbonate ($Na_2CO_3,10H_2O$). (It would be inconveniently moist, even semi-fluid, if a part of the carbonate had not previously been rendered anhydrous.) The *Sodii Bicarbonas*,

Fig. 17.



Preparation of bicarbonate of sodium.

U. S. P., is the commercial bicarbonate purified from any carbonate or traces of other salts by introducing it into a percolator and passing water through it till the washings cease to precipitate a solution of sulphate of magnesium, when the bicarbonate of sodium is removed from the percolator and dried on bibulous paper in a warm place.

A *crystal* of carbonate of sodium is carbonate of sodium plus water; on heating it, more or less of the water is evolved, and *anhydrous* carbonate of sodium is partially or wholly produced (*Sodæ Carbonas Exsiccata*, B. P., *Sodii Carbonas Exsiccata*, U. S. P.).

$$\underset{\text{Crystallized carbonate of sodium.}}{Na_2CO_3,10H_2O} \quad - \quad \underset{\text{Water.}}{10H_2O} \quad = \quad \underset{\text{Dried carbonate of sodium.}}{Na_2CO_3}$$

*Note on Nomenclature.*—*Anhydrous* bodies (from α, *a*, and ὕδωρ, *hudōr*, i. e. without water) are compounds from which water has been taken, but whose essential chemical properties are unaltered. Salts containing water are *hydrous* bodies; of these the larger portion are crystalline, and their water is then termed *water of crystallization*. Non-crystalline *hydrous* compounds were formerly spoken of as *hydrated* substances; *hydrates* are, however, a distinct class of bodies, salts derived from water by one-half of its hydrogen becoming displaced by an equivalent quantity of another radical. *Anhydrides* form still another distinct class of chemical substances; they are derived from acids; in short they are acids from which, not exactly water as water, but the elements of water have been removed, the essential chemical (acid) properties being thereby greatly altered. (For illustrations, see Index, "Anhydrides.")

*Water of Crystallization.*—The water in crystallized carbonate of sodium is in the solid condition, and, like ice and other fusible substances, requires heat for its liquefaction. Many salts (freezing-mixtures), when dissolved in water, give a very cold solution. This is because they and their solid water, if they have any, are then, absorbing some heat from surrounding media, converted into liquids. Take away from water some of its heat, the result is ice. Give to ice (at 32° F.) more heat than it contains already, the result is water (still at 32° F.). (Heat thus taken into a substance without increasing its temperature is said to become *latent*—from *latens*, hiding; it is no longer discoverable by the sense of touch or the thermometer. The term *latent* gives a somewhat incorrect idea, however, of the process; for our knowledge of the extent and readiness with which one form of force is convertible into another renders highly probable the assumption that heat is in these cases converted into motion, the latter enabling the molecules of a solid to take up the new positions demanded by their liquid condition.) The only apparent difference between ice and the water in such crystals as carbonate of sodium is that ice is solid water in the free, and water of crystallization solid water in the combined state. The former can only exist at and below freezing, the latter may exist at ordinary temperatures. Many salts,

however, which unite with little or even no water of crystallization at common temperatures, take up much, according to Guthrie, at very low temperatures, and such salts he calls *cryohydrates* (κρυος, *kruos*, icy cold, frost). On the other hand, all water of crystallization is dispelled at high temperatures. In chemical formulæ, the symbols representing water are usually separated by a comma from those representing salts. The crystals of acetate of sodium (of the third reaction) contain water in this loose state of combination—water of crystallization ($NaC_2H_3O_2,3H_2O$).

"*Soda-water.*"—A solution of bicarbonate of sodium in water charged with carbonic acid gas under pressure constitutes the official *Liquor Sodæ Effervescens*, B. P., and, like the "potash-water" of the shops, is a true medicine, an antacid. Ordinary "soda-water," however, is in many cases simply a solution of carbonic acid gas in water, and would be more appropriately termed "aërated water": any medicinal effect it may possess is due to the sedative influence of its carbonic acid gas on the coats of the stomach. At common temperatures water dissolves about its own volume of carbonic acid gas, both being under the same pressure. One pint of the official soda-water contains 30 grains of bicarbonate of sodium and a pint of carbonic acid gas; but the solution is under a pressure of seven atmospheres, so that seven pints of the gas at ordinary atmospheric pressure are required for the quantity mentioned.

*Solubility of Gases in Water.*—Whatever the weight and volume of a gas dissolved by a liquid at ordinary atmospheric pressure, that weight is doubled by double pressure, the two volumes of gas thereby being reduced to one, trebled at treble pressure, the three volumes of gas being reduced to one, quadrupled at quadruple pressure, the four volumes of gas being reduced to one, and so on. This is a general law (Henry and Dalton) regarding the solubility of gases in liquids under given temperatures. An average bottle of "soda-water" contains about four times the weight of carbonic acid gas which can exist in it without artificial pressure, so that on removing its cork three times its bulk escapes, its own bulk remaining dissolved.

*Note.*—Bicarbonate of sodium may also be medicinally administered in the form of lozenge (*Trochisci Sodii Bicarbonatis*, U. S. P.).

## Tartrate of Potassium and Sodium.

*Fifth Synthetical Reaction.*—To some hot strong solution of carbonate of sodium (about three parts) in a test-tube or larger vessel add acid tartrate of potassium (about four parts), till no more effervescence occurs; when the solution is cold, crystals of the tartrate of potassium and sodium (*Soda Tartarata*, B. P., *Potassii et Sodii Tartras*, U. S. P.), the old *Rochelle Salt*, will be deposited—($KNaC_4H_4O_6,4H_2O$). The crystals are usually halves of right rhombic prisms.

| $Na_2CO_3$ | + | $2KHC_4H_4O_6$ | = | $2KNaC_4H_4O_6$ | + | $H_2O$ | + | $CO_2$ |
|---|---|---|---|---|---|---|---|---|
| Carbonate of sodium. | | Acid tartrate of potassium. | | Tartrate of potassium and sodium. | | Water. | | Carbonic acid gas. |

FORMULÆ OF TARTRATES.

| | |
|---|---|
| Tartaric acid . . . . . | $HH\ C_4H_4O_6$ |
| Acid tartrate of potassium . . . . | $KH\ C_4H_4O_6$ |
| Tartrate of potassium and sodium . . | $KNaC_4H_4O_6$ |

Very close analogy will be noticed in the constitution of the molecules of these salts. When the other tartrates come under notice, it will be found they also have a similar constitution.

## Hypochlorite of Sodium.

*Sixth Synthetical Reaction.*—Pass chlorine (*vide* page 27) into a solution of carbonate of sodium. The result is a bleaching and disinfecting liquid, which, when made of prescribed strength (12 ounces of carbonate in 36 of water, charged by the washed chlorine from 15 fluidounces of hydrochloric acid and 4 ounces of black oxide of manganese), is the Solution of Chlorinated Soda (*Liquor Sodæ Chloratæ*) of the British Pharmacopœia. It is said to contain chloride of sodium (NaCl) and hypochlorite of sodium (NaClO), with some bicarbonate of sodium.

$$\underset{\text{Blk. oxide of manganese.}}{MnO_2} + \underset{\text{Hydrochloric acid.}}{4HCl} = \underset{\text{Chloride of manganese.}}{MnCl_2} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Chlorine.}}{Cl_2}$$

$$\underset{\text{Carbonate of sodium.}}{Na_2CO_3} + \underset{\text{Chlorine.}}{Cl_2} = \underset{\text{Colorinated soda.}}{NaCl,NaClO} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

*Liquor Sodæ Chlorinatæ*, U. S. P., is made by decomposing solution of carbonate of sodium by solution of chlorinated lime, sp. gr. 1.045.

$$\underset{\text{Carbonate of sodium.}}{2Na_2CO_3} + \underset{\text{Chlorinated lime.}}{CaCl_2,Ca2ClO} = \underset{\text{Chlorinated soda.}}{2(NaCl,NaClO)} + \underset{\text{Carbonate of calcium.}}{2CaCO_3}$$

## Other Sodium Compounds.

*Synthetical Reactions* portraying the chemistry of the remaining official compounds (namely, nitrate, sulphate, hyposulphite, borate, arseniate, and valerianate of sodium) are deferred until the several acidulous radicals of these salts have been described.

*Phosphate of Sodium.*—The preparation and composition of this salt will be most usefully studied after bone-ash, the source of it and other phosphates, has been described. Bone-ash is phosphate of calcium (see page 110).

*The Citro-Tartrate* (*Sodæ Citro-tartras Effervescens*, B. P.) is a mixture of bicarbonate of sodium (17 parts), citric acid (6), and tartaric acid (8), heated (to 200° or 220° F.) until the particles aggregate to a granular condition. When required for medicinal

use, a dose of the mixture is placed in water; escape of carbonic acid gas at once occurs, and an effervescing liquid results. This substance may be regarded as the official representative of the popular "Effervescing Citrate of Magnesia," which will be further noticed in connection with the salts of magnesium (page 120).

*Soda Powders* (*Pulveres Effervescentes*, U. S. P.) are formed of 30 grains of bicarbonate of sodium and 25 of tartaric acid wrapped separately in papers of different color. When mixed with water, tartrate of sodium results, a little bicarbonate also remaining.

*In the manufacture of Carbonate of Sodium* from chloride, the source of the sodium is chloride of sodium, and of the carbonic radical carbonate of calcium in the form of limestone. The chloride is first converted into sulphate, the sulphate is then roasted with coal and limestone, and the resulting *black-ash* lixiviated (*lixivia*, from *lix*, lye—water impregnated with alkaline salts: hence *lixiviation*, the operation of washing a mixture with the view of dissolving out salts). The lye, evaporated to dryness, yields crude carbonate of sodium (soda-ash). This process will be further described in connection with Carbonates.

---

*Deliquescence and Efflorescence.*—The carbonates of sodium and potassium, chemically closely allied, are readily distinguished physically. Carbonate of potassium quickly absorbs moisture from the air and becomes damp, wet, and finally fluid—it is *deliquescent* (*deliquescens*, melting away). Carbonate of sodium, on the other hand, yields some of its water of crystallization to the air, the crystals becoming white, opaque, and pulverulent—it is *effloresccent* (*efflorescens*, blossoming forth).

*Analogy of Sodium salts to Potassium salts.*—Other synthetical reactions might be described similar to those given under potassium, and thus citrate, iodide, bromide, iodate, bromate, chlorate, manganate, and permanganate of sodium, and many other salts be formed. But enough has been stated to show how analogous sodium is chemically to potassium. Such analogies will constantly present themselves. In few departments of knowledge are order and method more perceptible; in few is there as much natural law, as much *science*, as in chemistry.

*Substitution of Potassium and Sodium Salts for each other.*—Sodium salts being cheaper than potassium salts, the former may sometimes be economically substituted. That one is employed rather than the other, is often merely a result due to accident or fashion. But it must be borne in mind that in some cases a potassium salt will crystallize more readily than its sodium analogue, or that a sodium salt is stable when the corresponding potassium salt has a tendency to absorb moisture, or one may be more soluble than the other, or the two may have different medicinal effect. For these or similar reasons, a potassium salt has come to be used in medicine or trade, instead of the corresponding sodium salt, and *vice versâ*. Whenever the acidulous portion only is to be utilized, the least expensive salt of the class would nearly always be selected.

### (*b*) *Reactions having Analytical Interest.*

1. *The chief analytical reaction* for sodium is the *flame-test.* When brought into contact with a flame in the manner described under potassium (page 79), an intensely yellow color is communicated to the flame by any salt of sodium. This is highly characteristic—indeed, almost too delicate a test; for if the point of the wire be touched by the fingers, enough salt (which is contained in the moisture of the hand) adheres to the wire to communicate a very distinct sodium reaction. These statements should be experimentally verified, the chloride, sulphate, or any other salt of sodium being employed.

2. *Precipitant of Sodium.*—Sodium is the only metal whose common salts are all soluble in water. Hence no ordinary reagent can be added to a solution containing a sodium salt which shall give a precipitate containing the sodium A neutral or alkaline solution of a sodium salt gives, however, a granular precipitate of antimoniate of sodium ($Na_2H_2Sb_2O_7,6H_2O$) if well stirred or shaken with a solution of *antimoniate of potassium* ($K_2H_2Sb_2O_7$), but the reagent precipitates other metals, and is liable to decompose and become useless, and hence is seldom employed.

Antimoniate of potassium is made by adding, gradually, finely powdered metallic antimony to nitrate of potassium fused in a crucible so long as deflagration continues. The resulting mass is boiled with a large quantity of water, the solution filtered and preserved in a well-stoppered bottle; for the carbonic gas in the air is rapidly absorbed by the solution, antimonic acid being deposited.

3. Sodium salts, like those of potassium, are not volatile. Prove this fact by the means described when treating of the effect of heat on potassium salts (p. 79).

---

## QUESTIONS AND EXERCISES.

98. How is the official Solution of Soda prepared? Give a diagram or equation.

99. Explain the action of sodium or potassium on water. What colors do these elements respectively communicate to flame?

100. How much bicarbonate of sodium can be obtained from 2240 pounds of crystallized carbonate of sodium? Ans. 1316 lbs.

101. Acetate of Sodium: give formula, process, and equation.

102. Give a diagram showing the formation of Bicarbonate of Sodium.

103. Why is a mixture of dried and undried carbonate of sodium employed in the preparation of the bicarbonate?

104. State the difference between anhydrous and crystallized carbonate of sodium.

105. Define the terms *anhydrous*, *hydrous*, *hydrate*, *anhydride*.

106. What do you understand by *water of crystallization?*
107. What is the nature of "Soda-water?"
108. How many volumes of gas (reckoned as at ordinary atmospheric pressure) are contained in any given volume of the British official "Soda-water?"
109. What is the general law regarding the solubility of gases in liquids under pressure?
110. What is the systematic name of Rochelle salt, and how is the salt prepared?
111. What is the relation of Rochelle salt to cream of tartar and tartaric acid?
112. Give the mode of preparation and composition of Solution of Chlorinated Soda, aad express the process by a diagram.
113. How is the granular effervescing Citro-tartrate of Sodium prepared?
114. Define *Deliquescence*, *Efflorescence*, and *Lixiviation*.
115. What is the general relation of potassium salts to those of sodium?
116. How are sodium salts analytically distinguished from those of potassium?

---

## AMMONIUM.

Symbol $NH_4$ or Am. Atomic weight 18.

*Memoranda.*—The elements nitrogen and hydrogen, in the proportion of one atom to four ($NH_4$), are those characteristic of all the compounds about to be studied, just as potassium (K) and sodium (Na) are the characteristic elements of the potassium and sodium compounds. Ammonium is a univalent nucleus, root, or radical, like potassium or sodium; and the ammonium compounds closely resemble those of potassium or sodium. In short, if, for an instant, potassium or sodium be imagined to be compounds, the analogy between these three series of salts is complete. Ammonium is said to have been isolated by Weyl, as an unstable dark-blue liquid possessing a metallic lustre.

*Source.*—It is possible to produce ammonia from its elements, but the source of nearly all the ammoniacal salts met with in commerce is ammonia-gas ($NH_3$) obtained in distilling coals in the manufacture of ordinary illuminating gas. It is doubtless derived from the nitrogen of the plants from which the coal has been produced.

*Ammonia.*—When this gas ($NH_3$) comes into contact with water ($H_2O$), in the process of washing and cooling coal-gas, hydrate of ammonium ($NH_4HO$, or AmHO) is believed to be formed, the analogue of hydrate of potassium (KHO) or sodium (NaHO). The grounds for this belief are the observed analogy of the well-known ammoniacal salts to those of potassium and sodium, the similarity of action of solution of potash, soda, and ammonia on salts of most metals, and the existence of crystals of an analogous sulphur salt ($NH_4HS$).

*Chloride of Ammonium.*—The "ammoniacal liquor" of the gas-works is usually neutralized by hydrochloric acid, by which crude *chloride of ammonium* (sal-ammoniac) is produced.

$$NH_4HO + HCl = NH_4Cl + H_2O;$$

and from this salt, purified, the others used in pharmacy are directly or indirectly made. Chloride of ammonium (*Ammonii Chloridum*, B. P. and U. S. P.) occurs "in colorless, inodorous, translucent fibrous masses, tough, and difficult to powder, soluble in water [1 in 10 is the 'Solution of Chloride of Ammonium,' B. P.] and in rectified spirit." Chloride of ammonium generally contains slight traces of oxychloride of iron, tarry matter, and possibly chlorides of compound ammoniums (*vide* "Artificial Alkaloids" in Index).

*Sulphate of Ammonium*, $(NH_4)_2SO_4$, results when "ammoniacal liquor" is neutralized by oil of vitriol. It is largely used as a constituent of artificial manure in England, and when purified by recrystallization is employed in pharmacy (*Ammonii Sulphas*, U. S. P.).

*Volcanic Ammonia.*—The purest form of ammonia is that met with in volcanic districts, and obtained as a by-product in the manufacture of borax; the crude boracic acid as imported contains about 10 per cent. of ammonium salts, chiefly sulphate, and double sulphates of ammonium with magnesium, sodium, and manganese (Howard).

## Reactions having (*a*) General, (*b*) Synthetical, and (*c*) Analytical Interest.

### Ammonium-Amalgam. (?)

(*a*) *General Reaction.*—To forty or fifty grains of dry mercury in a *dry* test-tube, add one or two small pieces of sodium (freed from adhering naphtha by gentle pressure with a piece of filter-paper), and amalgamate by gently warming the tube. To this amalgam, when cold, add some fragments of chloride of ammonium and a strong solution of the same salt. The sodium amalgam soon begins to swell and rapidly increase in bulk, probably overflowing the tube. The light spongy mass produced is the so-called ammonium-amalgam, and the reaction is usually adduced as evidence of the existence of ammonium; the sodium of the amalgam unites with the chlorine of the chloride of ammonium, while the ammonium is supposed to form an amalgam with the mercury. As soon as formed the amalgam gives off hydrogen and ammonia gases; this decomposition is nearly complete after some minutes, and impure mercury remains.

### (*b*) *Reactions having Synthetical Interest.*

### Hydrate of Ammonium. Ammonia.

*First Synthetical Reaction.*—Heat a few grains of sal-ammoniac with about an equal weight of hydrate of calcium (slaked lime) dampened with a little water in a test-tube; ammonia gas is given off, and may be recognized by its well-known odor. It is very soluble in water. Pass a delivery-tube, fitted to the test-tube as described for the preparation of oxygen and hydrogen, into a second test-tube, at the bottom of which is a little water; again heat, the end of the delivery-tube being only just beneath the surface of the water (or, possibly, all the water might rush into the generating tubes, water absorbing ammonia gas with great avidity); solution of ammonia will be thus formed.

$$\underset{\text{Chloride of ammonium.}}{2NH_4Cl} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Ammonia gas.}}{2NH_3}$$

Ammonia gas is composed of one atom of nitrogen with three atoms of hydrogen; its formula is $NH_3$; two volumes of it contain one volume of nitrogen combined with three atoms or volumes of hydrogen. Its constituents have therefore in combining suffered condensation to one half their normal bulk. Its conversion into hydrate of ammonium may be thus shown:—

$$\underset{\text{Ammonia gas.}}{NH_3} + \underset{\text{Water.}}{H_2O} = \underset{\text{Hydrate of ammonium (ammonia).}}{NH_4HO \text{ or } AmHO}$$

*Solutions of Ammonia*, prepared by this process on a large scale and in suitable apparatus, are met with in pharmacy—the one (sp. gr. 0.891) containing 32.5 per cent., the other (sp. gr. 0.959), 10 per cent., by weight, of ammonia gas, $NH_3$, or 66.9 and 20.6 of ammonia, $NH_4HO$ (*Liquor Ammoniæ Fortior* and *Liquor Ammoniæ*, B. P. One part, *by measure*, of the former, and two of water form the latter). On the large scale, bottles are so arranged in a series as to condense all the ammonia evolved during the operation. *Aqua Ammoniæ Fortior*, U. S. P., sp. gr. 0.900, contains 26 per cent. of ammonia gas. *Aqua Ammoniæ*, U. S. P., has a sp. gr. of 0.960.

*Note.*—When water is mixed with strong solution of ammonia, expansion occurs. Thus, calculating from the stated specific gravities of the official solutions, 3000 parts by measure of the *Liquor Ammoniæ* weigh 2877 parts (959 × 3); whereas 1000 parts by measure of the *Liquor Ammoniæ Fortior* (weighing 891 parts) and 2000 measures of water (weighing 2000) weigh 2891 parts. Apparently 1000 volumes of the strong solution of ammonia with 2000 volumes of water yield not only 3000 volumes of mixture, but 14 parts by weight, or 15 volumes to spare, total 3015 volumes—an expansion of about half of one (.5) per cent.

## Acetate of Ammonium.

*Second Synthetical Reaction.*—To acetic acid and water in a test-tube, add powdered commercial carbonate (acid carbonate and carbamate) of ammonium till effervescence ceases; the resulting liquid, made of prescribed strength, is the official solution of Acetate of Ammonium ($NH_4C_2H_3O_2$) (*Liquor Ammoniæ Acetatis*, B. P., *Liquor Ammonii Acetatis*, U. S. P.).

$$\underset{\text{Acid carbonate and carbamate of ammonium.}}{(NH_4HCO_3)_2NH_4NH_2CO_2} + \underset{\text{Acetic acid.}}{4HC_2H_3O_2} = \underset{\text{Acetate of ammonium.}}{4NH_4C_2H_3O_2} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

Solution of acetate of ammonium could, of course, be just as easily made by reaction of acetic acid and solution of ammonia (AmHO), and, indeed, it was directed to be prepared in the British Pharmacopœia of 1864, but the liquid, owing to absence of dissolved carbonic acid, was considered to be too vapid for general use; hence the return to the older method.

## Carbonates of Ammonium.

*Commercial carbonate of ammonium* is made by heating a mixture of chalk and sal-ammoniac; chloride of calcium ($CaCl_2$) is produced, ammonia gas ($NH_3$) and water ($H_2O$) escape, and the ammoniacal carbonate distils, or rather sublimes,* in cakes (*Ammoniæ Carbonas*, B. P., *Ammonii Carbonas*, U. S. P.). The best form of apparatus to employ is a retort with a short wide neck and a cool receiver. On the large scale the retort is usually iron and the receiver earthenware or glass; on the small scale glass vessels are employed. The salt is purified by resublimation at a low temperature; 150° F. is said to be sufficient.

This salt, the empirical formula of which is $N_4H_{16}C_3O_8$, is probably a mixture of two molecules of acid carbonate or bicarbonate of ammonium ($2NH_4HCO_3$) and one of a salt termed carbamate of ammonium ($NH_4NH_2CO_2$). The latter belongs to an important class of salts known as carbamates, but it is the only one of interest to the pharmacist. Cold water extracts it from the commercial carbonate of ammonium, leaving the acid carbonate of ammonium undissolved, if the amount of liquid used be very small. In water, carbamate soon changes into neutral carbonate of ammonium,

$$NH_4NH_2CO_2 + H_2O = (NH_4)_2CO_3 \text{ or } Am_2CO_3;$$

so that an aqueous solution of commercial carbonate of ammonium contains both acid carbonate and neutral carbonate of ammonium.

---

* *Sublimation* (from *sublimis*, high). Vaporization of a solid substance by heat, and its condensation on an upper and cooler part of the vessel or apparatus in which the operation is performed.

If to such a solution some ordinary solution of ammonia be added, a solution of *neutral carbonate of ammonium* is obtained; and this is the common reagent always found on the shelves of the analytical laboratory.

$$AmHCO_3 + AmHO = Am_2CO_3 + H_2O.$$

Neutral carbonate of ammonium is the salt formed on adding strong solution of ammonia to the commercial carbonate in preparing a pungent mixture for toilet smelling-bottles; but it is unstable, and on continued exposure to air is reduced to a mass of crystals of the acid carbonate or bicarbonate of ammonium. Bicarbonate of ammonium ($NH_4HCO_3$) is also produced on passing carbonic acid gas into an aqueous solution of commercial carbonate.

According to Divers, the sublimed product of the first distillation of chalk and sal-ammoniac is a mixture of carbamate and carbonate of ammonium, the latter losing some ammonia gas on redistillation, and carbamate with bicarbonate forming the resulting commercial salt. Dr. Divers considers that the salt now met with in trade contains one molecule of acid carbonate to one of carbamate, and not two of the former to one of the latter, as was the case when the analyses were made from which the foregoing formulæ were deduced. Statements by manufacturers and analyses by the author point to the same conclusion.

*Sal Volatile* (*Spiritus Ammoniæ Aromaticus*, B. P. and U. S. P.) is a spirituous solution of ammonia (AmHO), neutral carbonate of ammonium ($Am_2CO_3$), and the oils of nutmeg and lemon (and lavender, U. S. P.). Fetid spirit of ammonia (*Spiritus Ammoniæ Fœtidus*, B. P.) is an alcoholic solution of the volatile oil of assafœtida mixed with solution of ammonia. "Solution of Carbonate of Ammonia," B. P., is formed by dissolving half an ounce of the salt in ten ounces of water. *Spiritus Ammoniæ*, U. S. P., is an alcoholic solution of ammonia.

## Nitrate of Ammonium.

*Third Synthetical Reaction.*—To some diluted nitric acid add carbonate of ammonium, until, after well stirring, a slight ammoniacal odor remains. The solution contains Nitrate of Ammonium (*Ammoniæ Nitras*, B. P., *Ammonii Nitras*, U. S. P.).

$$\underset{\text{Acid carbonate and carbamate of ammonium.}}{(NH_4HCO_3)_2,\ NH_4NH_2CO_2} + \underset{\text{Nitric acid.}}{4HNO_3} = \underset{\text{Nitrate of ammonium.}}{4NH_4NO_3} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Carbonic acid gas.}}{3CO_2}.$$

From a strong hot solution of nitrate of ammonium crystals may be obtained containing much water ($NH_4NO_3, 12H_2O$). On heating these to about 310° F. water escapes, and the anhydrous salt remaining ($NH_4NO_3$), may be poured on to an iron plate. On further heating the powdered nitrate, it is resolved into *nitrous oxide* gas (the so-called laughing gas) and water.

$$NH_4NO_3 = N_2O + 2H_2O.$$

Nitrous oxide is thus prepared for use as an anæsthetic. When required for inhalation, it should be washed from any possible trace of acid or nitric oxide, by being passed through solution of potash, and through solution of ferrous sulphate.

Nitrous oxide is slightly soluble in warm water, more so in cold. It supports combustion almost as well as oxygen. By pressure it may be liquefied to a colorless fluid, and by simultaneous cooling solidified.

## Citrate, Phosphate, and Benzoate of Ammonium.

*Fourth Synthetical Reaction.*—To solution of citric acid ($H_3C_6H_5O_7$ or $H_3\overline{Ci}$) add solution of ammonia (AmHO) until the well-stirred liquid smells faintly of ammonia; the product is Solution of Citrate of Ammonium ($Am_3\overline{Ci}$) (*Liquor Ammoniæ Citratis*, B. P.).

*Phosphate of Ammonium* ($Am_2HPO_4$) (*Ammoniæ Phosphas*, B. P.) and *Benzoate of Ammonium* ($AmC_7H_5O_2$) (*Ammoniæ Benzoas*, B. P., *Ammonii Benzoas*, U. S. P.) are also made by adding solution of ammonia to phosphoric acid ($H_3PO_4$) and benzoic acid ($HC_7H_5O_2$) respectively, evaporating (keeping the ammonia in slight excess by adding more of its solution), and setting aside for crystals to form. The official Solution of Acetate of Ammonium could be made in the same way; but, when prepared with Carbonate of Ammonium, the liquid remains charged with carbonic acid, and has a less vapid flavor.

| | | | | | | |
|---|---|---|---|---|---|---|
| $H_3C_6H_5O_7$ Citric acid. | + | 3AmHO Ammonia. | = | $Am_3C_6H_5O_7$ Citrate of ammonium. | + | $3H_2O$ Water. |
| $H_3PO_4$ Phosphoric acid. | + | 2AmHO Ammonia. | = | $Am_2HPO_4$ Phosphate of ammonium. | + | $2H_2O$ Water. |
| $HC_7H_5O_2$ Benzoic acid. | + | AmHO Ammonia. | = | $AmC_7H_5O_2$ Benzoate of ammonium. | + | $H_2O$ Water. |

Phosphate of ammonium occurs in transparent colorless prisms, soluble in water, insoluble in spirit; benzoate in crystalline plates, soluble in water and in spirit.

*Ammonii Iodidum*, U. S. P., is made by decomposing the two bodies iodide of potassium and sulphate of ammonium, which give iodide of ammonium and sulphate of potassium; the latter salt is separated by adding alcohol to the cooled solution, when, by reason of its insolubility in alcohol, it crystallizes out, and the separated solution of iodide of ammonium is then evaporated to dryness.

*Bromide of Ammonium* (*Ammonii Bromidum*, B. P., and U. S. P.) will be noticed in connection with Hydrobromic Acid and other Bromides.

## Oxalate of Ammonium.

*Fifth Synthetical Reaction.*—To a nearly boiling solution of 1 part of oxalic acid in about 8 of water add carbonate of ammonium until the liquid is neutral to *test-paper* (see following paragraph), filter while hot, and set aside for crystals ($(NH_4)_2C_2O_4.H_2O$) to form. The mother-liquor is useful as a reagent in analysis; 1 of the salt in 40 of water constitutes "Solution of Oxalate of Ammonia," B. P.

$$\underset{\text{Oxalic acid.}}{2H_2C_2O_4} + \underset{\text{Carbonate of ammonium.}}{N_4H_{16}C_2O_6} = \underset{\text{Oxalate of ammonium.}}{2(NH_4)_2C_2O_4} + \underset{\text{Carbonic acid gas.}}{3CO_2} + \underset{\text{Water.}}{2H_2O}$$

*Neutralization.*—Thus far, in reactions, the student has avoided excess of either acid matter on the one hand, or alkaline matter on the other, by the rough aid of taste, cessation of effervescence, presence or absence of odor, etc. More delicate aid is afforded by *test-papers.*

*Test-papers.*—*Litmus* (B. P.) is a blue vegetable pigment, prepared from various species of *Roccella* lichen, exceedingly sensitive to the action of acids, which turn it red. When thus reddened, alkalies (potash, soda, and ammonia) and other soluble hydrates readily turn it blue. The student should here test for himself the delicacy of this action by experiments with paper soaked in solution of litmus and dipped into very dilute solutions of acids, acid salts ($KHC_4H_4O_6$, *e. g.*), alkalies, and such neutral salts as nitrate of potassium, sulphate of sodium, or chloride of ammonium.

*Tincture of Litmus* (B. P.).—1 ounce of litmus is macerated for two days in 10 fluidounces of proof spirit, and the solution poured off from insoluble matter.

*Blue litmus paper* (B. P.) is "unsized white paper steeped in tincture of litmus, and dried by exposure to the air." *Red litmus paper* (B. P.) is "unsized white paper steeped in tincture of litmus which has been previously reddened by the addition of a very minute quantity of sulphuric acid, and dried by exposure to the air."

*Turmeric paper*, similarly prepared from tincture of turmeric (1 of turmeric root or rhizome to 6 of rectified spirit macerated for seven days), is occasionally useful as a test for alkalies, which turn its yellow to brown; acids do not affect it.

## Sulphydrate of Ammonium.

*Sixth Synthetical Reaction.*—Pass sulphuretted hydrogen gas ($H_2S$) through a small quantity of solution of ammonia in a test-tube, until a portion of the liquid no longer causes a white precipitate in solution of sulphate of magnesium (Epsom salt); the product is solution of sulphydrate (or sulphide) of ammonium ($NH_4HS$), the "Solution of Sulphide of Ammo-

nium" of the British Pharmacopœia, a most valuable chemical reagent, as will presently be apparent.

$$NH_4HO + H_2S = NH_4HS + H_2O.$$

"*Solution of Sulphide of Ammonium*," B. P., is made by passing the gas prepared in the apparatus described below into 3 fluidounces of solution of ammonia (*Liquor Ammoniæ*) so long as the gas continues to be absorbed, then adding 2 more ounces of solution of ammonia, and preserving the solution in a well-stoppered bottle.

*Sulphuretted hydrogen* is a compound of noxious odor; hence the above operation, and many others, described further on, in which this gas is indispensable, can only be performed in the open air, or in a fume-cupboard, a chamber so contrived that deleterious gases and vapors shall escape into a chimney in connection with the external air. In the above experiment, the small quantity of gas required can be made in a test-tube, after the manner of hydrogen itself. To two or three fragments of sulphide of iron (FeS), add water and then sulphuric acid; the gas is at once evolved, and may be conducted by a tube into the solution of ammonia. Sulphate of iron remains dissolved in the water.

$$FeS + H_2SO_4 = H_2S + FeSO_4.$$

*Crystals of sulphydrate of ammonium* ($NH_4HS$) may be obtained on bringing ammonia gas ($NH_3$) and sulphuretted hydrogen ($H_2S$) together at a low temperature. They are soluble in water without decomposition.

*Sulphuretted-Hydrogen Apparatus.*—As no heat is necessary in making sulphuretted hydrogen, the test-tube of the foregoing operation may be advantageously replaced by a bottle, especially when larger quantities of the gas are required. In analytical operations, the gas should be purified by passing it through water contained in a second bottle.

The most convenient arrangement for experimental use is prepared as follows: Two common wide-mouth bottles are selected, the one having a capacity of about half a pint, the other a quarter pint; the former may be called the generating-bottle, the latter the wash-bottle. Fit two corks to the bottles. Through each cork bore two holes by a round file or other instrument, of such a size that glass tubing of about the diameter of a quill pen shall fit them tightly. Through one of the holes in the cork of the generating-bottle pass a funnel-tube, so that its extremity may nearly reach the bottom of the bottle. To the other hole adapt a piece of tubing, 6 inches long, and bent in the middle to a right angle. A similar "elbow-tube"

is fitted to one of the holes in the cork of the wash-bottle, and another elbow-tube, one arm of which is long enough to reach to near the bottom of the wash-bottle, fitted to the other hole. Removing the corks, two or three ounces of water are now

Fig. 18.

Sulphuretted-Hydrogen Apparatus.

poured into each bottle, an ounce or two of sulphide of iron put into the generating-bottle, and the corks replaced. The elbow-tube of the generating-bottle is now attached by a short piece of India-rubber tubing to the long-armed elbow-tube of the wash-bottle, so that gas coming from the generator may pass through the water in the wash-bottle. The delivery-tube of the wash-bottle is then lengthened by attaching to it, by India-rubber tubing, another piece of glass tubing several inches in length. The apparatus is now ready for use. Strong sulphuric acid is poured down the funnel-tube in small quantities at a time, until brisk effervescence is established, and more added from time to time as the evolution of gas becomes slow. The gas passes through the tubes into the wash-bottle, where, as it bubbles up through the water, any trace of sulphuric acid, or other matter mechanically carried over, is arrested, and thence flows out at the delivery-tube into any vessel or liquid that may be placed there to receive it. The generator must be occasionally dismounted, and the sulphate of iron washed out.

*Luting* (*lutum*, mud).—If the corks of the above apparatus are sound, and tube-holes well made, no escape of gas will occur. If rough corks have been employed, or the holes are not cylindrical, linseed-meal lute may be rubbed over the defective parts. The lute is prepared by mixing linseed-meal with water to the consistence of dough. A neat appearance may be given to the lute by gently rubbing a well-wetted finger over its surface.

### (c) *Reactions having Analytical Interest (Tests).*

*First Analytical Reaction.*—To a solution of any salt of ammonium (the chloride, for example) in a test-tube, add solution of caustic soda (or solution of potash, or a little slaked lime); ammonia gas is at once evolved, recognized by its well-known odor.

$$NH_4Cl + NaHO = NH_3 + H_2O + NaCl.$$

Though ammonium itself cannot be kept in the free state, its compounds are stable. Ammonia is easily expelled from these compounds by action of the stronger alkalies, caustic potash, soda, or lime. As a matter of exercise, the student should here draw out equations in which acetate ($NH_4C_2H_3O_2$), sulphate ($Am_2SO_4$), nitrate ($NH_4NO_3$) or any other ammoniacal salt not already having the odor of ammonia, is supposed to be under examination; also representing the use of the other hydrates, potash (KHO) or slaked lime (Ca2HO).

The *odor* of ammonia gas is perhaps the best means of recognizing its presence; but the following tests are also occasionally useful. Into the test-tube in which the ammonia gas is evolved insert a glass rod moistened with hydrochloric acid (that is, with the solution of hydrochloric acid gas, conveniently termed hydrochloric acid, the *Acidum Hydrochloricum* of the Pharmacopœias); white fumes of chloride of ammonium will be produced.

$$NH_3 + HCl = NH_4Cl.$$

Hold a piece of moistened red litmus paper in a tube in which ammonia gas is present; the red color will be changed to blue.

*Second Analytical Reaction.*—To a few drops of a solution of an ammonium salt add a drop or two of hydrochloric acid and a like small quantity of solution of perchloride of platinum ($PtCl_4$); a yellow crystalline precipitate of the double chloride of platinum and ammonium ($PtCl_4 2NH_4Cl$) will be produced, similar in appearance to the corresponding salt of potassium, the remarks concerning which (p. 78) are equally applicable to the precipitate under notice.

*Third Analytical Reaction.*—To a moderately strong solution of an ammonium salt add a strong solution of tartaric acid, and shake or well stir the mixture; a white granular precipitate of acid tartrate of ammonium will be formed.

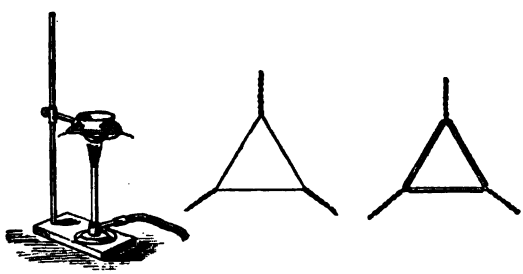
For data from which to draw out an equation representing this action, see the remarks and formulæ under the analogous salt of potassium (p. 78).

*Fourth Analytical Fact.*—Evaporate a few drops of a solution of an ammonium salt to dryness, or place a fragment of a

salt in the solid state on a piece of platinum foil, and heat in a flame; the salt is readily *volatilized.* As already noticed, the salts of potassium and sodium are *fixed* under these circumstances, a point of difference of which advantage will frequently be taken in analysis. A porcelain crucible may often be advantageously substituted for platinum foil in experiments on volatilization.

Salts of ammonium with the more complex acidulous radicals seldom volatilize unchanged when heated. The oxalate, when warmed, loses its water of crystallization, and at a higher temperature decomposes, yielding carbonic oxide, carbonic acid gas, ammonia gas, water (the three latter sometimes in combination), and several organic substances. The phosphate yields more or less phosphoric acid as a residue.

Fig. 19. Fig. 20. Fig. 21.



Triangular Supports for Crucibles.

*A wire-triangle* may be used in supporting crucibles. It is made by twisting together each pair of ends of three (5 or 6 inch) crossed pieces of wire (Fig. 20). A piece of tobacco-pipe stem (about 2 inches) is sometimes placed in the centre of each wire before twisting, the transference of any metallic matter to the sides of the crucible being thus prevented (Fig. 21.)

### *Practical Analysis.*

With regard to those experiments which are useful rather as means of detecting the presence of potassium, sodium, and ammonium, than as illustrating the preparation of salts, the student should proceed to apply them to certain solutions of any of the salts of potassium, sodium, and ammonium, with the view of ascertaining which metal is present; that is, proceed to practical analysis.* A little thought

* Such solutions are prepared in educational laboratories by a tutor. They should, under other circumstances, be mixed by a friend,

will enable him to apply these reactions in the most suitable order and to the best advantage for the contemplated purpose: but the following arrangements are perhaps as good as can be devised:—

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE METALS, POTASSIUM, SODIUM, AMMONIUM.

Add caustic soda to a small portion of the solution to be examined, and warm the mixture in a test-tube; the odor of ammonia gas at once reveals the presence of an ammonium salt.

If ammonium be not present, apply the perchloride-of-platinum test; a yellow precipitate proves the presence of potassium.

(It will be observed that potassium can only be detected in the absence of ammonium, salts of the latter radical giving similar precipitates.)

The flame-test is sufficient for the recognition of sodium.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE, TWO, OR ALL THREE OF THE ALKALI METALS.

Commence by testing a small portion of the solution for an ammonium salt. If present, make a memorandum to that effect, and then proceed to get rid of the ammoniacal compound to make way for the detection of potassium: advantage is here taken of the volatility of ammonium salts and the fixity of those of potassium and sodium. Evaporate the original solution to dryness in a small basin, transfer the solid residue to a porcelain crucible, and heat the latter to a low redness, or until dense white fumes (of ammoniacal salts) cease to escape. (See Fig. 19.) This operation should be conducted in a fume-cupboard,

---

as it is not desirable to know previously what is contained in the substance about to be analyzed.

The analysis of solutions containing only one salt serves to impress the memory with the characteristic tests for the various metals and other radicals, and familiarize the mind with chemical principles. Medical students seldom have time to go further than this. More thorough analytical and general chemical knowledge is only acquired by working on such mixtures of bodies as are met with in actual practice, beginning with solutions which may contain any or all the members of a group. Hence in this manual, two Tables of short directions for analyzing are given under each group. Pharmaceutical students should follow the second Table.

to avoid contamination of the air of the apartment. When the crucible is cold, dissolve out the solid residue with a small quantity of hot water, and test the solution for potassium by the perchloride-of-platinum test, and for sodium by the flame-test.

If ammonium is proved to be absent, the original solution may, of course, be at once tested for potassium and sodium.

*Flame-test.*—The violet tint imparted to flame by potassium salts may be seen when masked by the intense yellow color due to sodium, if the flame be observed through a piece of dark-blue glass, a medium which absorbs the yellow rays of light.

*Note on Nomenclature.*—The operations of *evaporation* and heating to redness, or *ignition*, are frequently necessary in analysis, and are usually conducted in the above manner. If vegetable or animal matter be also present, carbon is set free, and ignition is accompanied by *carbonization;* the material is said to *char*. When all carbonaceous matter is burnt off, the crucible being slightly inclined and its cover removed to facilitate combustion, and mineral matter, or *ash*, alone remains, the operation of *incineration* has been effected.

*Note on the Classification of Elements.*—The compounds of potassium, sodium, and ammonium have many analogies. Their carbonates, phosphates, and other common salts are soluble in water. The atoms of the radicals themselves are univalent—that is, displace or are displaced by one atom of hydrogen. In fact, they constitute by their similarity in properties a distinct group or family. All the elements thus naturally fall into classes—a fact that should constantly be borne in mind, and evidence of which should always be sought. It would be impossible for the memory to retain the details of chemistry without a system of classification and leading principles. Classification is also an important feature in the art as well as in the science of chemistry; for without it practical analysis could not be undertaken. The classification adopted in this volume is founded, as far as possible, on the quantivalence of the elements, but chiefly on their analytical relations.

---

## QUESTIONS AND EXERCISES.

117. Why are ammoniacal salts classed with those of potassium and sodium?

118. Mention the sources of the ammonium salts.

119. Describe the appearance and other characters of Chloride of Ammonium.

120. Give the formula of Sulphate of Ammonium.

121. Adduce evidence of the existence of ammonium.

122. How are the official Solutions of Ammonia prepared? Give diagrams.

123. How is the official Solution of Acetate of Ammonium prepared?

124. What is the composition of commercial Carbonate of Ammonium?

125. Define *sublimation*.

126. What ammoniacal salts are contained in *Spiritus Ammoniæ Aromaticus*?

127. Give diagrams or equations illustrating the formation of Citrate, Phosphate, and Benzoate of Ammonium.

128. Give the formula of Oxalate of Ammonium.

129. Show how hydrate of ammonium may be converted into sulphydrate.

130. Describe the preparation of Sulphuretted Hydrogen gas.

131. Enumerate and explain the tests for ammonium.

132. How is potassium detected in a solution in which ammonium has been found?

133. Give equations illustrating the action of hydrate of sodium on acetate of ammonium; hydrate of potassium on sulphate of ammonium; and hydrate of calcium on nitrate of ammonium.

134. What are the effects of acids and alkalies on litmus and turmeric?

135. Describe the analysis of an aqueous liquid containing salts of potassium, sodium, and ammonium.

136. What meanings are commonly assigned to the terms *evaporation*, *ignition*, *carbonization*, and *incineration*?

137. Write a short article descriptive of the analogies of potassium, sodium, and ammonium, and their compounds.

---

## BARIUM, CALCIUM, MAGNESIUM.

These three elements have many analogies. Their atoms are bivalent.

### BARIUM.

Symbol Ba. Atomic weight 137.

The analytical reactions only of this metal are of interest to the general student of pharmacy. The nitrate ($Ba2NO_3$) and chloride ($BaCl_2,2H_2O$) are the soluble salts in common use in analysis (Chloride of Barium, B. P., and "Solution of Chloride of Barium," 1 in 10 of water, B. P.); and these and others are made by dissolving the native carbonate ($BaCO_3$), *Barii Carbonas*, U. S. P., the mineral *witherite*, in acids, or by heating the other common natural compound of barium, the sulphate, *heavy white* or *heavy spar* ($BaSO_4$), with coal, which yields sulphide of barium (BaS),

$$BaSO_4 + C_4 = 4CO + BaS,$$

and dissolving the sulphide in appropriate acids. When the nitrate is strongly heated, it is decomposed, the oxide of barium or *baryta*

(BaO) remaining. Baryta, on being moistened, assimilates the elements of water with great avidity, and yields hydrate of barium (Ba2HO). The latter is tolerably soluble, giving *baryta-water;* and from this solution crystals of hydrate of barium are obtained on evaporation.

The operations above described may all be performed in test-tubes and small porcelain crucibles heated by the gas-flame. Quantities of 1 oz. to 1 lb. require a coke-furnace.

*Peroxide of barium* ($BaO_2$) is formed on passing air over baryta heated to low redness. By the action of dilute hydrochloric acid it yields solution of *peroxide of hydrogen* ($H_2O_2$), the old *oxygenated water*.

*Quantivalence.*—The atom of barium is bivalent, Ba″.

## Reactions having Analytical Interest (Tests).

*First Analytical Reaction.*—To the solution of any soluble salt of barium (nitrate or chloride, for example) add dilute sulphûric acid; a white precipitate is obtained. Set the test-tube aside for two or three minutes, and when some of the precipitate has fallen to the bottom pour away most of the supernatant liquid, add strong nitric acid, and boil; the precipitate is insoluble.

The production of a white precipitate by sulphuric acid, insoluble even in hot nitric acid, is highly characteristic of barium. The name of this precipitate is sulphate of barium; its formula is $BaSO_4$.

*Antidotes.*—In cases of poisoning by soluble barium salts, obvious antidotes would be solutions of alum or of any sulphates, such as those of magnesium and sodium (Epsom salt, Glauber's salt).

*Second Analytical Reaction.*—To a barium solution add solution of the yellow chromate of potassium ($K_2CrO_4$); a pale yellow precipitate ($BaCrO_4$) falls. Add acetic acid to a portion of the chromate of barium; it is insoluble. Add hydrochloric or nitric acid to another portion; it is soluble.

"*Neutral Chromate.*"—The red chromate (or bichromate) of potassium ($K_2CrO_4,CrO_3$) must not be used in this reaction, or the barium will be only imperfectly precipitated; for the red salt gives rise to the formation of free acid, in which chromate of barium is to some extent soluble:—

$$K_2CrO_4,CrO_3 + 2BaCl_2 + H_2O = 2BaCrO_4 + 2KCl + 2HCl.$$

Yellow chromate is obtained on adding carbonate of potassium, in small quantities at a time, to a hot solution of the red chromate until effervescence ceases; a little more red chromate is then added to insure decomposition of any slight excess of carbonate of potassium.

$$K_2CrO_4,CrO_3 + K_2CO_3 = 2K_2CrO_4 + CO_2.$$

For analytical purposes, solution of a neutral chromate is still more readily prepared by simply adding solution of ammonia to solution of red chromate of potassium, until the liquid turns yellow, and, after stirring, smells of ammonia.

$$K_2CrO_4,CrO_3 + 2NH_4HO = 2KNH_4CrO_4 + H_2O.$$

*Other Analytical Reactions.*—To a barium solution add a soluble carbonate (carbonate of ammonium ($Am_2CO_3$) will generally be rather more useful than others); a white precipitate of carbonate of barium ($BaCO_3$) results.——To more of the solution add an alkaline phosphate or arseniate (phosphate of sodium ($Na_2HPO_4$) is the most common of these chemically analogous salts, but phosphate of ammonium ($Am_2HPO_4$) or arseniate ($Am_2HAsO_4$) will subsequently have the preference); white phosphate of barium ($BaHPO_4$), insoluble in pure water but slightly soluble in aqueous solutions of some salts, or arseniate of barium ($BaHAsO_4$), both soluble even in acetic and other weak acids, are precipitated.——To another portion add oxalate of ammonium ($Am_2C_2O_4$); white oxalate of barium ($BaC_2O_4$) is precipitated, soluble in the diluted mineral acids, and sparingly so in acetic acid.——The silico-fluoride of barium ($BaSiF_6$) is insoluble, and falls readily if an equal volume of spirit of wine be added to the solution under examination after the addition of hydrofluosilicic acid ($H_2SiF_6$).——Barium salts, moistened with hydrochloric acid, impart a greenish color to flame.

*Mem.*—Good practice will be found in writing out equations descriptive of each of the foregoing reactions.

---

## QUESTIONS AND EXERCISES.

138. What is the quantivalence of barium?

139. Write down the formulæ of oxide, hydrate, chloride, nitrate, carbonate, and sulphate of barium; and state how these salts are prepared.

140. Describe the preparation of peroxide of hydrogen.

141. Which of the tests for barium are most characteristic? Give an equation of the reactions.

142. Name the antidote in cases of poisoning by soluble barium salts, and explain its action.

# CALCIUM.

Symbol Ca. Atomic weight 40.

Calcium compounds form a large proportion of the crust of our earth. Carbonate of calcium is met with as chalk, marble, limestone, calc-spar, etc., the sulphate, as gypsum or plaster of Paris ("Plaster of Paris, native sulphate of calcium — $CaSO_4$, $2H_2O$— deprived of water by heat."—B. P.), and alabaster, the silicate in many minerals, the fluoride of calcium as fluor-spar. The phosphate is also a common mineral. The element itself is only isolated with great difficulty. The atom of calcium is bivalent, $Ca''$.

## Reactions having Synthetical Interest.

### Chloride of Calcium.

*First Synthetical Reaction.*—To some hydrochloric acid add carbonate of calcium (chalk, or, the purer form, white marble, *Marmor Album*, B. P. and U. S. P.) ($CaCO_3$) until effervescence ceases, filter; solution of chloride of calcium ($CaCl_2$), the most common soluble salt of calcium, is formed.

$$\underset{\text{Carbonate of calcium.}}{CaCO_3} + \underset{\text{Hydrochloric acid.}}{2HCl} = \underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

This solution contains carbonic acid, and will give a precipitate of carbonate of calcium on the addition of lime-water. It may be obtained quite neutral by well boiling before filtering off the excess of marble. It is a serviceable test-liquid in analytical operations.

Solution of chloride of calcium evaporated to a syrupy consistence readily yields crystals. These are extremely deliquescent. The solution, evaporated to dryness, and the white residue strongly heated, gives solid anhydrous chloride of calcium in a porous form. The resulting agglutinated lumps (*Calcii Chloridum*, B. P. and U. S. P.) are much used for drying gases, and for freezing certain liquids from water. The salt is soluble in alcohol. One part in ten of water constitutes a useful test liquid, "Solution of Chloride of Calcium," B. P. Four parts in five of water form the "Solution (saturated) of Chloride of Calcium," B. P.

*Mem.*—The practical student has already met with solution of chloride of calcium as a by-product or secondary product in the preparation of carbonic acid gas.

Marble often contains ferrous carbonate ($FeCO_3$), which in the above process becomes converted into ferrous chloride, rendering the chloride of calcium impure:—

$$\underset{\text{Ferrous carbonate.}}{FeCO_3} + \underset{\text{Hydrochloric acid.}}{2HCl} = \underset{\text{Ferrous chloride.}}{FeCl_2} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

If absolutely pure chloride of calcium be required, a few drops of the solution should be poured into a test-tube or test-glass, diluted with water, and examined for iron (by adding sulphydrate of ammonium, which gives a black precipitate with salts of iron), and, if the latter is present, hypochlorite of calcium (in the form of chlorinated lime) and slaked lime should be added to the remaining bulk of the liquid, and the whole boiled for a few minutes, whereby iron (as ferric hydrate) is thus precipitated; on filtering, a pure solution of chloride of calcium is obtained:—

$$\underset{\text{Ferrous chloride.}}{4FeCl_2} + \underset{\text{Hypochlorite of calcium.}}{Ca2ClO} + \underset{\text{Hydrate of calcium.}}{4CaH_2O_2} + \underset{\text{Water.}}{2H_2O}$$
$$= \underset{\text{Ferric hydrate.}}{2(Fe_26HO)} + \underset{\text{Chloride of calcium.}}{5CaCl_2}$$

This is the official process, and may be imitated on the small scale after adding a minute piece of iron to a fragment of the marble before dissolving in acid.

The names, formulæ, and reactions of these compounds of iron will be best understood when that metal comes under treatment.

## Oxide of Calcium (Quick Lime).

*Second Synthetical Reaction.*—Place a small piece of chalk in a strong grate-fire or furnace and heat until a trial fragment, chipped off from time to time and cooled, no longer effervesces on the addition of acid; caustic lime, CaO (*Calx*, B. P. and U. S. P.), remains.

$$\underset{\text{Carbonate of calcium (chalk).}}{CaCO_3} = \underset{\text{Oxide of calcium (lime).}}{CaO} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

*Note.*—Etymologically considered, this action is analytical (ἀναλύω, *analuo*, I resolve) and not synthetical (σύνθεσις, *sūnthesis*, a putting together); but conventionally it is synthetical, and not analytical; for in this, the usual sense, and the sense in which the words are used throughout this book, synthesis is the application of chemical action with the view of producing something, analysis the application of chemical action with the view of finding out the composition of a substance. In the etymological view of the matter, there is scarcely an operation performed either by the analyst or by the manufacturer but includes both analysis and synthesis—that is, includes interchange or *metathesis*.

*Lime-kilns.*—On the large scale the above operation is carried on in what are termed *lime-kilns* (*Kiln*, Saxon, *cyln*, from *cylene*, a furnace).

## Hydrate of Calcium (Slaked Lime).

*Slaked Lime.*—When cold, add to the lime about half its weight of water, and notice the evolution of steam and other evidence of strong action; the product is *slaked* lime or hydrate of calcium (Ca2HO) (*Calcis Hydras*, B. P.), with whatever slight natural impurities the lime might contain. The slaking of hard or "stony" lime may be accelerated by using hot water.

$$\underset{\text{Lime.}}{CaO} + \underset{\text{Water.}}{H_2O} = \underset{\substack{\text{Hydrate of calcium}\\ \text{(slaked lime).}}}{Ca2HO}$$

*Lime-water.*—Place the hydrate of calcium in about a hundred times its weight of water; in a short time a saturated solution, known as *lime-water* (*Liquor Calcis*, B. P. and U. S. P.), results. It contains about 16 grains of hydrate of calcium (Ca2HO), equivalent to about 11 or 12 grains of lime (CaO), in one (Imperial) pint.

*Strong Solution of Lime.*—Slaked lime is much more soluble in aqueous solution of sugar than in pure water. The *Liquor Calcis Saccharatus*, B. P., is such a solution, containing 2 ounces of sugar and 188 grains of hydrate of calcium (Ca2HO), equivalent to 142 grains of lime (CaO), in 1 (Imp.) pint. It is a more efficient precipitant of hydrates and carbonates than lime-water. The official process is as follows: Mix 1 ounce of lime and 2 of sugar by trituration in a mortar. Transfer the mixture to a bottle containing 1 pint of water, and, having closed this with a cork, shake it occasionally for a few hours. Finally separate the clear solution with a siphon and keep it in a stoppered bottle.

Solutions of hydrate of calcium absorb carbonic acid gas on exposure to air, a semicrystalline precipitate of carbonate being deposited. When the saccharated solution is heated, there is precipitated a compound consisting of three molecules of lime with one of sugar.

## Carbonate of Calcium.

*Third Synthetical Reaction.*—To a solution of chloride of calcium add excess of carbonate of sodium, or about 5 parts of dry chloride to 13 of carbonate; a white precipitate of carbonate of calcium (*Calcis Carbonas Præcipitata*, B. P., *Calcii Carbonas Præcipitata*, U. S. P.) ($CaCO_3$) results. If the solutions of the salts be made hot before admixture, and the whole set aside for a short time, the particles aggregate to a greater extent than when cold water is used, and the product

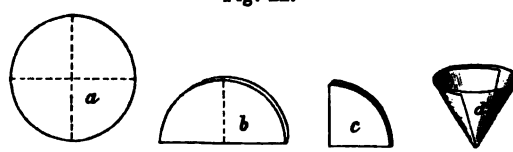
is finely granular or slightly crystalline. The official variety is thus prepared.

$$\underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Carbonate of sodium.}}{Na_2CO_3} = \underset{\text{Carbonate of calcium.}}{CaCO_3} + \underset{\text{Chloride of sodium.}}{2NaCl}$$

Collect and purify this *Precipitated Chalk* by pouring the mixture into a paper cone supported by a funnel, and, when the liquid has passed through the filter, pour water over the precipitate three or four times until the whole of the chloride of sodium is washed away. This operation is termed *washing a precipitate.* When dried by aid of water-bath (p. 111) or other means, the precipitate is fit for use.

*Filtering-paper*, or *bibulous-paper* (from *bibo*, to drink), is simply good unsized paper made from the best white rags—white blotting-paper, in fact, of unusually good quality. Students' or analysts' filters, on which to collect precipitates, are circular pieces (*a*) of this paper, from three to six inches in diameter, twice folded (*b*, *c*), and then opened out so as to form a hollow cone (*d*). Square pieces are rounded by scissors *after* folding. The cone is supported by a glass or earthenware funnel.

Fig. 22.




Construction of Paper Filters.

Filters should *always* be cut round so as to form a cone. If the square piece of paper is folded without being so cut or trimmed, an ugly angular filter results, from which it is difficult to wash all "mother-liquor" (the solution of chloride of sodium is the "mother-liquor" in the previous reaction). If a spirituous or other volatile fluid is being passed through such an angular filter, much of the liquid will also be wasted by evaporation from the unnecessarily large surface exposed.

*Washing-bottle.*—Precipitates are best washed by a fine jet of water directed on to the different parts of the filter. A common narrow-necked bottle of about half-pint capacity (Fig. 23) is fitted with a cork; two holes are bored through the cork, the one for a glass tube reaching to the bottom of the bottle within, and externally bent to a slightly acute angle, the other for a tube bent to a slightly obtuse angle, the inner arm terminating just within the bottle. The outer arms may be about 3 inches in length. The extremity of the outer arm continuous with the long tube should be previously drawn out to a fine capillary opening by holding the

original tube, before cutting, in a flame, and, when soft, slowly pulling the halves away from each other until the heated portion is reduced to the thinness of a knitting-needle. The tube is now cut at the thin part by a file, and the sharp edges rounded off by placing in

Fig. 23. Fig. 24.

Washing-bottles.

a flame for a second or two. The outer extremity of the shorter tube should also be made smooth in the flame. The apparatus being put together, and the bottle nearly filled with water, air, blown through the short tube by the lungs, forces water out in a fine stream at the capillary orifice.

*A hot-water washing flask* (Fig. 24) is made in the manner just described, the tubes and cork being fitted into a flask instead of a bottle.

*Decantation.*—Precipitates may also be washed by allowing them to settle, pouring off the supernatant liquid (Fig. 25), agitating with

Fig. 25. Fig. 26. Fig. 27.

Decantation. Decantation. Siphon in action.

water, again allowing to settle, and so on. This is washing by *decantation* (*de*, from, *canthus*, an edge). If a stream of liquid flowing from a basin or other vessel exhibits any tendency to run down

the outer side of the vessel, it should be guided by a glass rod placed against the point whence the stream emerges (Fig. 26).

If the vessel be too large to handle with convenience, the wash-water may be drawn off by a *siphon*, as shown in miniature in Fig. 27. A siphon is a tube of glass, metal, gutta-percha, or India-rubber, bent into the form of a V or U, filled with water, and inverted; one end immersed in the wash-water, and the other allowed to hang over the side of the vessel; so long as the outer orifice of the instrument is below the level of the liquid in the vessel, so long will that liquid flow from within outwards until the vessel be empty.*

*Prepared* carbonate of calcium (*Creta Præparata*, B. P. and U. S. P.) is merely washed chalk (*Creta*, B. P. and U. S. P.) or *whiting*, only that in Pharmacy fashion demands that the chalk be in little conical lumps, about the size of thimbles, instead of in the larger rolls characteristic of "whiting." Wet whiting pushed, portion by portion, through a funnel, and each separately dried, gives the conventional *Creta Præparata*. Its powder is amorphous.

*Testa Præparata*, U. S. P., is powdered oyster-shell, similarly treated. It is an inferior kind of prepared chalk.

## Phosphate of Calcium.

*Fourth Synthetical Reaction.*—Digest bone-ash (bones burnt in an open crucible with free access of air till all animal and carbonaceous matter has been removed—impure phosphate of calcium (*Os Ustum*, B. P.)) with nearly twice its weight of hydrochloric acid (diluted with three or four times its bulk of water), in a test-tube or larger vessel; the phosphate is dissolved.

$$\underset{\text{Phosphate of calcium (impure).}}{Ca_3 2PO_4} + \underset{\text{Hydrochloric acid.}}{4HCl} = \underset{\text{Acid phosphate of calcium.}}{CaH_4 2PO_4} + \underset{\text{Chloride of calcium.}}{2CaCl_2}$$

Dilute with water, filter, boil, and when cold add excess of solution of ammonia; the phosphate of calcium, now pure (*Calcis Phosphas*, B.P.; *Calcii Phosphas Præcipitata*, U.S.P.), is reprecipitated as a light white amorphous powder. After well washing, the precipitate should be dried over a *water-bath*

* *The nature of the action of a siphon* is simple. The column of water in the outer limb is longer, and therefore heavier, than the column of similar area in the inner limb. (The length of the inner limb must be reckoned from the surface of the liquid, the portion below the surface playing no part in the operation.) Being heavier, it naturally falls by gravitation, the liquid in the shorter limb instantly following, because pressed upwards by the air. The air, be it observed, exerts a similar amount of pressure on the liquid in the outer limb: in short, atmospheric pressure causes the retention of liquid in the instrument, while gravitation determines the direction of the flow.

(see below), or at a temperature not exceeding 212°, to prevent undue aggregation of the particles.

$$\underset{\text{Acid phosphate of calcium.}}{CaH_4P_2O_8} + \underset{\text{Chloride of calcium.}}{2CaCl_2} + \underset{\text{Ammonia.}}{4AmHO} = \underset{\text{Phosphate of calcium (pure).}}{Ca_2P_2O_8} + \underset{\text{Chloride of ammonium.}}{4AmCl} + \underset{\text{Water.}}{4H_2O}$$

Bone-ash or bone-earth contains small quantities of carbonate and sulphide of calcium. These are decomposed in the above process by the acid, chloride of calcium being formed; on boiling the mixture, carbonic acid gas and sulphuretted hydrogen gas are evolved. Any carbonaceous or siliceous matter, etc., is removed by filtration. In bones the phosphate of calcium is always accompanied by a small quantity of an allied substance, phosphate of magnesium; a trace of fluoride of calcium ($CaF_2$) is also present.

A *Water-bath* for the evaporation of liquids or for drying moist solids at temperatures below 212° F. is an iron, tin, or earthenware pan, the mouth of which can be narrowed by iron or tin diaphragms of various sizes, and having orifices adapted to the diameters of evaporating dishes or plates. In the British Pharmacopœia, "when a *water-bath* is directed to be used, it is to be understood that this term refers to an apparatus by means of which water or its vapor, at a temperature not exceeding 212°, is applied to the outer surface of a vessel containing the substance to be heated, which substance may thus be subjected to a heat near to, but necessarily below, that of 212°. In the *steam-bath* the vapor of water at a temperature above 212°, but not exceeding 230°, is similarly applied." Evaporation *in vacuo* is performed by simply placing the vessel of liquid over or by the side of a small reservoir of strong sulphuric acid, or other absorbent of moisture, on the plate of an air-pump, covering with a capacious glass hood or "receiver," and exhausting.

*Bone-black*, or *Animal Charcoal* (*Carbo Animalis*, B. P. and U. S. P), is the residue obtained on subjecting dried bones (*Os*, U. S. P.) to a red heat without access of air. The operation may be imitated by heating a few fragments of bone in a covered porcelain crucible in a fume-chamber until smoke and vapor cease to be evolved. Purified Animal Charcoal (*Carbo Animalis Purificatus*, B. P. and U. S. P.) is obtained by digesting animal charcoal (16 parts) in hydrochloric acid (10 parts) and water (20 parts) in a warm place for a day or two, filtering, thoroughly washing, drying over a water-bath, and igniting the product in a closely covered crucible. The reaction

is the same as that just described; that is to say, the acid removes the phosphate of calcium from the carbon of animal charcoal by forming soluble acid phosphate and chloride of calcium.

*Wood Charcoal* (*Carbo Ligni*, B. P. and U. S. P.) is wood similarly ignited without access of air.

*Decolorizing power of Animal Charcoal.*—Animal charcoal, in small fragments, is the material employed in decolorizing solutions of common brown sugar with the view of producing white lump sugar. Its power and the nearly equal power of an equivalent quantity of the purified variety may be demonstrated on solution of litmus or logwood.

*Phosphate of Sodium.*—Phosphate of calcium is converted into phosphate of sodium (*Sodæ Phosphas*, B. P., *Sodii Phosphas*, U. S. P.) ($Na_2HPO_4,12H_2O$) as follows: Mix, in a mortar, 3 ounces of ground bone-earth with 1 fluidounce of sulphuric acid; set aside for twenty-four hours to promote reaction; mix in about 3 ounces of water, and put in a warm place for two days, a little water being added to make up for that lost by evaporation; stir in another 3 ounces of water, warm the whole for a short time, filter, and wash the residual sulphate of calcium on the filter to remove adhering acid phosphate of calcium; concentrate the *filtrate* (the liquid portion), which is a solution of acid phosphate of calcium, to about 3 ounces, filter again, if necessary, add solution of (about $4\frac{1}{2}$ ounces of crystals of) carbonate of sodium to the hot filtrate until a precipitate (a phosphate of calcium, $CaHPO_4$) ceases to form, and the fluid is faintly alkaline; filter, evaporate, and set aside to crystallize.

Phosphate of sodium occurs "in transparent colorless rhombic prisms, terminated by four converging planes, efflorescent, tasting like common salt." One part to ten of water constitutes "Solution of Phosphate of Soda," B. P. This is an official as well as the ordinary process. The following equations show the two decompositions which occur during the operations:—

| | | | | | | |
|---|---|---|---|---|---|---|
| $Ca_3 2PO_4$ Phosphate of calcium. | + | $2H_2SO_4$ Sulphuric acid. | = | $CaH_4 2PO_4$ Acid phosphate of calcium | + | $2CaSO_4$ Sulphate of calcium. |
| $CaH_4 2PO_4$ Acid phosphate of calcium. | + | $Na_2CO_3$ Carbonate of sodium. | = | $Na_2HPO_4$ Phosphate of sodium. | + | $H_2O$ Water. |
| | + | $CO_2$ Carbonic acid gas. | + | $CaHPO_4$ Phosphate of calcium. | | |

Ordinary phosphate of sodium ($Na_2HPO_4,12H_2O$) effloresces rapidly in the air until nearly half its water has escaped, when

it has a permanent composition represented by the formula $Na_2HPO_4,7H_2O$. Phosphate of sodium has an alkaline reaction, the usual effect of acids on it being the removal of half its sodium and formation of the salt $NaH_2PO_4,H_2O$.

## Hypochlorite of Calcium.

*Fifth Synthetical Reaction.*—Pass chlorine, generated as already described, into damp slaked lime contained in a piece of wide tubing, open at the opposite end to that in which the delivery-tube is fixed. (A test-tube, the bottom of which has been accidentally broken, is very convenient for such operations.) The product is ordinary *bleaching-powder*, a compound of hypochlorite and chloride of calcium, commonly called *chloride of lime* (*Calx Chlorata*, B. P., *Calx Chlorinata*, U.S.P.).

| $MnO_2$ | + | $4HCl$ | = | $MnCl_2$ | + | $2H_2O$ | + | $Cl_2$ |
|---|---|---|---|---|---|---|---|---|
| Black oxide of manganese. | | Hydrochloric acid. | | Chloride of manganese. | | Water. | | Chlorine. |

| $2CaH_2O_2$ | + | $2Cl_2$ | = | $2H_2O$ | + | $CaCl_2O_2$, | $CaCl_2$ |
|---|---|---|---|---|---|---|---|
| Hydrate of calcium. | | Chlorine. | | Water. | | Hypochlorite of calcium. | Chloride of calcium. |

Chlorinated lime, exposed to air and moisture, as in disinfecting the air of sick-rooms, slowly yields hypochlorous acid ($HClO$). Free hypochlorous acid soon breaks up into water, chloric acid ($HClO_3$), and free chlorine. Chloric acid is also unstable, decomposing into oxygen, water, chlorine, and perchloric acid ($HClO_4$). The small quantity of hypochlorous acid, diffused through an apartment when bleaching-powder is exposed thus, yields fourteen-fifteenths of its chlorine in the form of chlorine gas—one of the most efficient of known disinfectants.

*Constitution of Bleaching-powder.*—Treated with alcohol, bleaching-powder does not yield its chloride of calcium to the solvent, hence the powder is not a mere mixture of chloride and hypochlorite of calcium: water, also, does not dissolve out first one salt, and then the other, but both together, in the molecular proportions of the above formula. On the other hand, when the aqueous solution is cooled, or evaporated *in vacuo*, crystals are obtained which Kingzett has shown to be nearly pure hypochlorite of calcium, the solution containing chloride of calcium. While the former fact indicates that the powder is a compound, and not a mere mixture, the latter indicates that it is a feeble compound; an adhesion of molecules of hypochlorite and chloride, as shown in the equation, rather than any more intimate or closer combination of atoms.

If it be regarded as a single rather than a double salt then the following formula may be employed, $Ca \begin{cases} Cl \\ ClO \end{cases}$.

*Bleaching-liquor.*—Digest chlorinated lime in water, in which the bleaching compound is soluble, filter from the undissolved lime, and test the bleaching powers of the clear liquid by adding a few drops to a decoction of logwood slightly acidulated. One pound of this bleaching-powder, shaken several times during three hours with 1 gallon of water, forms Solution of Chlorinated Lime (*Liquor Calcis Chloratæ*, B. P.).

## Gummate of Calcium.

*Gummate of Calcium* is the only official calcium salt that remains to be noticed. This compound is, in short, *arabin*, the ordinary Gum Acacia, or Gum Arabic (*Acaciæ Gummi*, B. P. and U. S. P.), a substance too well known to need description. A solution of gum arabic in water (*Mucilago Acaciæ*, B. P. and U. S. P.) yields a white precipitate of oxalate of calcium on the addition of solution of oxalate of ammonium. Or a piece of gum burnt to an ash in a porcelain crucible yields a calcareous residue, which, dissolved in dilute acids, affords characteristic reactions with any of the following analytical reagents for calcium. In some specimens of gum arabic a portion of the calcium is displaced by an equivalent quantity of potassium or magnesium. The gummic radical may be precipitated as opaque gelatinous gummate of lead by the addition of solution of oxyacetate of lead (*Liquor Plumbi Subacetatis*, B. P. and U. S. P.) to an aqueous solution of gum. These statements may be experimentally verified by the practical student.

*Tragacanth* (*Tragacantha*, B. P. and U. S. P.) is a mixture of soluble arabinoid gum and a variety of calcium gum insoluble in water, termed *bassorin*. With water a gelatinous mucilage is formed (*Mucilago Tragacanthæ*, B. P. and U. S. P.).

### Reactions having Analytical Interest (Tests).

*First Analytical Reaction.*—Add sulphuric acid, highly diluted, to a calcium solution contained in a test-tube, or small test-glass; sulphate of calcium ($CaSO_4,2H_2O$) is formed, but is *not* precipitated, it being, unlike sulphate of barium, slightly soluble in water.

*Solution of Sulphate of Calcium.*—A quarter of an ounce of that (dried) form of sulphate of calcium known as plaster of Paris ($CaSO_4$)

digested in one pint of water for a short time, with occasional shaking, and the mixture filtered, yields the official test-liquid termed "Solution of Sulphate of Lime," B. P. About 400 parts of the solution contain 1 of calcium.

*Second Analytical Reaction.*—Add yellow chromate of potassium ($K_2CrO_4$) to a calcium solution slightly acidified with acetic acid; chromate of calcium ($CaCrO_4$) is probably formed, but it is *not* precipitated. Barium is precipitated by the chromic radical.

*These two negative reactions* are most valuable in analysis, as every precipitant of calcium is also a precipitant of barium; but the above two reagents are precipitants of barium only. Hence calcium which when alone can be readily detected by the following reactions, cannot by any reaction be detected in the presence of barium. But by the sulphuric or chromic test barium is easily removed, and then either of the following reagents will throw down the calcium.

*Other Analytical Reactions.*—Add carbonate of ammonium, phosphate of sodium, arseniate of ammonium, and oxalate of ammonium to calcium solutions as described under the analytical reactions of barium, and write out descriptive equations. The precipitates correspond in appearance to those of barium; their constitution is also similar, hence their correct formulæ can easily be deduced. Of these precipitants oxalate of ammonium is that most commonly used as a reagent for calcium salts, barium being absent. The oxalate of calcium is insoluble in acetic, but soluble in hydrochloric or nitric acids.——Calcium compounds impart a reddish color to the flame.

---

## QUESTIONS AND EXERCISES.

143. Enumerate some of the common natural compounds of calcium.

144. Explain, by an equation, the action of hydrochloric acid on marble. What official compounds result?

145. Why is chloride of calcium used as a desiccator for gases?

146. How would you purify Chloride of Calcium which has been made from ferruginous marble? Give diagrams.

147. Write a few lines on the chemistry of the lime-kiln.

148. In what sense is the conversion of chalk into lime an analytical action?

149. What occurs when lime is slaked?

150. To what extent is lime soluble in water? to what in syrup?

151. Describe the preparation of the official Precipitate of Carbonate of Calcium; in what does it differ from Prepared Chalk?

152. In what does filtering paper differ from other kinds of paper?

153. Explain the construction of "a washing bottle" for cleansing precipitates by water.

154. Describe *decantation*.

155. Describe the construction and manner of employment of a siphon.

156. Explain the mode of action of a siphon.

157. What is the difference between Bone, Bone-earth, and Precipitated Phosphate of Calcium?

158. How is "Bone-earth" purified for use in medicine?

159. Explain the action of hydrochloric acid on Animal Charcoal in the conversion of *Carbo Animalis* into *Carbo Animalis Purificatus*.

160. What is the chemical difference between *Carbo Animalis* and *Carbo Ligni*?

161. Give equations showing the conversion of Phosphate of Calcium into Phosphate of Sodium.

162. Write a short article on the manufacture, composition, and uses of "bleaching-powder."

163. How may calcium be detected in Gum Arabic?

164. State the chemical nature of Tragacanth.

165. To what extent is sulphate of calcium soluble in water?

166. Can calcium be precipitated from an aqueous solution containing barium?

167. Barium being absent, what reagents may be used for the detection of calcium? Which is the chief test?

---

## MAGNESIUM.

Symbol Mg. Atomic weight 24.

*Source.*—Magnesium is abundant in nature in the form of magnesian or mountain limestone, or *dolomite*, a double carbonate of magnesium and calcium in common use as a building-stone (*e. g.*, the Houses of Parliament and the School of Mines in London), and *magnesite*, a tolerably pure carbonate of magnesium, though too "stony" for direct use in medicine, even if very finely powdered. Chloride of magnesium and sulphate of magnesium (Epsom salt) also occur in sea-water and in the water of many springs. A monohydrous sulphate ($MgSO_4,H_2O$) termed *kieserit* occurs near Stassfurt in Prussia. Metallic magnesium may be obtained from the chloride by the action of sodium. It burns readily in the air, emitting a dazzling light due to the white heat to which the resulting particles of magnesia (MgO) are exposed. The chloride employed as a source of the metal is obtained by dissolving the carbonate in hydrochloric acid, adding some chloride of ammonium, evaporating to dryness, heating the residue in a flask (on the small scale a large test-tube or Florence flask) until the chloride of ammonium is all volatilized and the chloride of magnesium remains as a clear fused liquid. The latter is poured on to a clean earthenware slab. The chloride of ammonium

prevents reaction between chloride of magnesium and water in the last stages of the operation and consequent formation of oxide (or oxychloride) of magnesium and hydrochloric acid gas.

*Quantivalence.*—The atom of magnesium is bivalent, Mg″.

## Reactions having Synthetical Interest.

### Sulphate of Magnesium.

*First Synthetical Reaction.*—To a few drops of sulphuric acid and a little water in a test-tube, made hot (or to larger quantities in larger vessels), add powdered native carbonate of magnesium, *magnesite*, $MgCO_3$, until effervescence ceases, subsequently boiling to aid in the expulsion of the carbonic acid gas. The filtered liquid is a solution of sulphate of magnesium ($MgSO_4$), crystals of which, *Epsom salt* ($MgSO_4,7H_2O$) (*Magnesiæ Sulphas*, B. P., *Magnesii Sulphas*, U. S. P.), may be obtained on evaporating most of the water, and setting the concentrated solution aside to cool. This is an ordinary manufacturing process. Instead of magnesite, *dolomite*, the common magnesian limestone (carbonate of magnesium and calcium, $CaCO_3,MgCO_3$) may be employed, any iron being removed by evaporating the solution (filtered from the sulphate of calcium produced) to dryness, gently igniting to decompose sulphate of iron, dissolving in water, filtering from oxide of iron, and crystallizing. (If neither mineral be at hand, the practical student may use a little of the ordinary manufactured carbonate of pharmacy, for the chemical action is almost identical, and it is the chemistry and not, just now, the commercial economy of the matter that he is studying. The manufacturer must, of course, commence with one of the above mineral carbonates furnished by nature, from that make his sulphate, and from the latter, as will be seen directly, make the pure pulverulent carbonate of pharmacy.)

$$\underset{\text{Carbonate of magnesium.}}{MgCO_3} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Sulphate of magnesium.}}{MgSO_4} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

Sulphate of magnesium readily crystallizes in large, colorless, transparent, rhombic prisms; but, from concentrated solutions, the crystals are deposited in short thin needles, a form more convenient for manipulation, solution, and general use in medicine.

Iron may be detected in sulphate of magnesium by adding the common alkaline solution of chlorinated lime or chlorinated soda to some aqueous solution of the salt; brown hydrate of iron ($Fe_2 6HO$) being precipitated. Sulphydrate ammonium will also give a black precipitate if iron be present.

## Carbonates of Magnesium.

*Second Synthetical Reaction.*—To solution of sulphate of magnesium add solution of carbonate of sodium and boil; the resulting precipitate is *light* carbonate of magnesium (*Magnesiæ Carbonas Levis*, B. P., *Magnesii Carbonas*, U. S. P.), a white, partly amorphous, partly minutely crystalline mixture of carbonate and hydrate of magnesium ($3MgCO_3,Mg2HO,4H_2O$). A denser, slightly granular precipitate of similar chemical composition (*Magnesiæ Carbonas*, B. P.) is obtained on mixing strong solutions of the above salts, evaporating to dryness, then removing the sulphate of sodium by digesting the residue in hot water, filtering, washing, and drying the precipitate.

$$\underset{\text{Sulphate of magnesium.}}{4MgSO_4} + \underset{\text{Carbonate of sodium.}}{4Na_2CO_3} + \underset{\text{Water.}}{H_2O} = \underset{\text{Official carbonate of magnesium.}}{3MgCO_3,Mg2HO} + \underset{\text{Sulphate of sodium.}}{4Na_2SO_4} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

The official proportions for the light carbonate are 10 of sulphate of magnesium and 12 of crystals of carbonate of sodium, each dissolved in 80 of cold water, the solutions mixed, boiled for 15 minutes, the precipitate collected on a filter, well washed, drained, and dried over a water-bath. The heavier carbonate is made with the same proportions of salts, each dissolved in 20 instead of 80 of water, the mixture evaporated quite to dryness, and the residue washed by decantation or filtration until all sulphate of sodium is removed (shown by a white precipitate—sulphate of barium—ceasing to form on the addition of solution of chloride or nitrate of barium to a little of the filtrate.)

*Another* (*Pattinson's*) *Process.*—Considerable quantities of carbonate of magnesium are now prepared by treating dolomite (see p. 117) with carbonic acid under pressure. Of the two carbonates the magnesian is dissolved first, and is precipitated from the clear liquid by the heat of a current of steam. (See next reaction.)

*Third Synthetical Reaction.*—Pass carbonic acid gas, generated as described on page 71, into a mixture of water and carbonate of magnesium contained in a test-tube. After some time, separate undissolved carbonate by filtration; the filtrate contains carbonate of magnesium dissolved by carbonic acid. When of a strength of about 13 grains in one ounce, such a solution constitutes "*Fluid Magnesia*" (*Liquor Magnesiæ Carbonatis*, B. P.).

Officially, 1 pint is directed to be made from freshly prepared carbonate. The latter is obtained by adding a hot solution of 2 ounces

of sulphate of magnesium in half a pint of water to one of 2½ ounces of crystals of carbonate of sodium in another half pint of water, boiling the mixture for a short time (to complete decomposition), filtering, thoroughly washing the precipitate, placing the latter in 1 pint of distilled water, and transmitting carbonic acid gas through the liquid (say, at the rate of three or four bubbles per second) for an hour or two, then leaving the solution in contact with the gas under slight pressure for twenty-four hours, and, finally, filtering from undissolved carbonate, and, after passing in a little more gas, keeping in a well-corked bottle. Slight pressure is best created by placing the carbonate and water in a bottle fitted with a cork and tubes as for a wash-bottle (p. 96 or 108), conveying the gas by a tube which reaches to the bottom, and allowing excess of gas to flow out by the upper tube, the external end of which is continued to the bottom of a common phial containing about an inch of mercury. The phial should be loosely plugged with cotton-wool, to prevent loss of metal by spurting during the flow of the gas through it. (Each inch in depth of mercury through which the gas escapes corresponds to about half a pound pressure on every square inch of surface within the apparatus.)

Heat a portion of the solution; true carbonate of magnesium containing combined water ($MgCO_3,3H_2O$) is precipitated. The water in this compound is probably in the state of water of crystallization, for a salt having the same composition is deposited in crystals by the spontaneous evaporation of the solution of carbonate of magnesium. The official "carbonate" ($3MgCO_3,Mg2HO,4H_2O$) is another of these very common *hydrous* compounds.

Exposed to cold, the solution of "fluid magnesia" sometimes affords large thick crystals ($MgCO_3,5H_2O$), which, in contact with the air, lose water, become opaque, and then have the composition of those deposited by evaporation ($MgCO_3,3H_2O$).

*Liquor Magnesii Citratis*, U. S. P., is a solution of carbonate of magnesium and bicarbonate of potassium in excess of citric acid and syrup of citric acid.

## Oxide of Magnesium (Magnesia).

*Fourth Synthetical Reaction.*—Heat light dry carbonate of magnesium in a porcelain crucible over a lamp (or in a larger earthen crucible in a furnace) till it ceases to effervesce on adding, to a small portion, water and acid; the residue is light magnesia (MgO) (*Magnesia Levis*, B. P.). The same operation on the heavy carbonate yields heavy magnesia (MgO) (*Magnesia*, B. P.). Both are sometimes spoken of as "calcined magnesia" (*Magnesia*, U. S. P.). A given weight of the official light magnesia occupies three and a half times the bulk of the weight of heavy magnesia.

| $3MgCO_3,Mg2HO$ | = | $4MgO$ | + | $H_2O$ | + | $3CO_2$ |
|---|---|---|---|---|---|---|
| Official carbonate of magnesium. | | Oxide of magnesium. | | Water. | | Carbonic acid gas. |

A trace only of magnesia is dissolved by pure water. Moisten a grain or two of magnesia with water, and place the paste on a piece of red litmus paper; the wet spot, after a time, becomes blue, showing that the magnesia is slightly soluble.

"*Effervescing Citrate of Magnesia*," so-called, is generally a mixture of bicarbonate of sodium, citric acid, tartaric acid, sugar, either carbonate or sulphate of magnesium, or both, and flavoring essences. True citrate of magnesium is easily made by heating together calcined magnesia and citric acid; it is frequently prescribed in France in doses of two ounces.

The official Effervescing "Solution of Citrate of Magnesia" (*Liquor Magnesiæ Citratis*, B. P.) is made by dissolving carbonate of magnesium in slight excess of solution of citric acid, adding lemon-syrup, placing the diluted liquid in an aërated-water bottle, dropping in crystals of bicarbonate of potassium, corking, "wiring," and shaking till the crystals are dissolved. This preparation is similar to the *Liquor Magnesii Citratis* of the United States Pharmacopœia.

The formula of citrate of magnesium deposited from solution is $Mg_3 2C_6H_5O_7, 14H_2O$.

### Reactions having Analytical Interest (Tests).

*First Analytical Reaction.*—Add solution of hydrate or carbonate of ammonium to a magnesian solution (sulphate for example) and warm the mixture in a test-tube; the precipitation of part only of the magnesium as hydrate ($Mg2HO$) or carbonate ($MgCO_3$) occurs. Add now to a small portion of the mixture of precipitate and liquid a considerable excess of solution of chloride of ammonium; the precipitate is dissolved.

This is an important reaction, especially as regards carbonate of magnesium, the presence of chloride of ammonium enabling the analyst to throw out from a solution barium and calcium by an alkaline carbonate, magnesium being retained. The cause of this reaction is found in the tendency of magnesium to form soluble double salts with potassium, sodium, or ammonium. In analysis, the chloride of ammonium should be added before the carbonate, as it is easier to prevent precipitation than to redissolve a precipitate once formed.

*Second Analytical Reaction.*—To some of the solution resulting from the last reaction, add solution of phosphate of sodium or ammonium; phosphate of magnesium and ammonium ($MgNH_4PO_4$) is precipitated.——3*d*. To another portion add arseniate of ammonium; arseniate of magnesium and ammonium ($MgNH_4AsO_4$) is precipitated.

*Note.*—Barium and calcium are also precipitated by alkaline phosphates and arseniates. The other precipitants of magnesium are also precipitants of barium and calcium. In other words, there is no *direct* test for magnesium. Hence the analyst always removes any barium or calcium by an alkaline carbonate, as above indicated; the phosphate of sodium, or arseniate or phosphate of ammonium, then become very delicate tests of the presence of magnesium. In speaking of magnesium tests, the absence of barium and calcium salts is to be understood.

## QUESTIONS AND EXERCISES.

168. Name the natural sources of the various salts of magnesium.
169. Give a process for the preparation of Epsom salt.
170. Draw diagrams illustrative of the formation of sulphate of magnesium from *magnesite* and from *dolomite*.
171. Show by an equation the process for the preparation of the official Carbonate of Magnesium.
172. What circumstances determine the two different states of aggregation of the *Magnesiæ Carbonas* and *Magnesiæ Carbonas Levis*?
173. What are the relations of *Magnesia* and *Magnesia Levis* to the official Carbonates of Magnesium?
174. How much denser is the one than the other?
175. Is magnesia soluble in water?
176. How is "Fluid Magnesia" prepared?
177. Mention the effects of heat and cold on "Fluid Magnesia."
178. How much magnesia (MgO) can be obtained from 100 grains of Epsom salt?
179. Calculate the amount of official Carbonate of Magnesium which will yield 100 grains of magnesia.
180. Can magnesium be detected in presence of barium and calcium?
181. Describe the analysis of an aqueous liquid containing salts of barium, calcium, and magnesium.
182. How may magnesium be precipitated from solutions containing ammoniacal salts?

### *Quantivalence.*

On reviewing the foregoing statements regarding compounds of the three univalent radicals, potassium, sodium, and ammonium, and the three bivalent elements, barium, calcium, and magnesium, the doctrine of quantivalence will be more clearly understood, and its usefulness more apparent. Quantivalence, or the value of atoms, is, in short, in chemistry, closely allied to value in commercial barter. A number of articles, differing much in weight, appearance, and general characters, may be of equal money value; and if these be

regarded, for convenience, as having a sort of unit of value, others worth double as much might be termed bivalent, three times as much trivalent, and so on. In like manner, chemical radicals, no matter whether elementary, like potassium (K), iodine (I), or sulphur (S), or compound, like those of nitrates ($NO_3$), sulphates ($SO_4$), or acetates ($C_2H_3O_2$), have a given chemical value in relation to each other, and are exchangeable for, and will unite with each other to an extent determined by that value.

Most chemical salts apparently, though probably not really, have two parts, a basylous and an acidulous, the one quantivalently balancing the other. The formulæ of the chief of these radicals and their quantivalence are given below. Examples of formulæ of salts containing univalent, bivalent, and trivalent radicals are also appended.

QUANTIVALENCE OF COMMON RADICALS.

| Univalent Radicals, or *Monads*. | | Bivalent Radicals, or *Dyads*. | | Trivalent Radicals, or *Triads*. | |
|---|---|---|---|---|---|
| Acidulous. | Basylous. | Acidulous. | Basylous. | Acidulous. | Basylous. |
| H | H | O | Ca | $PO_4$ | As |
| Cl | K | $SO_4$ | Mg | $BO_3$ | Sb |
| I | Na | $CO_3$ | Zn | $C_6H_5O_7$ | Bi |
| HO | $NH_4$ | $C_2O_4$ | Cu | $AsO_3$ | $Fe^{III}$ (ic) |
| $NO_3$ | Ag | $C_4H_4O_6$ | Hg(ic) | $AsO_4$ | or |
| $C_2H_3O_2$ | Hg(ous) | S | Fe(ous) | $C_4H_3O_5$ | $Fe^{VI}_2$(ic) |

*Note.*—The hydrogen (H) in the basylous parts of salts has entirely different functions from the hydrogen (H) in the acidulous part. The latter gives compounds commonly termed *hydrides* (*e. g.*, $AsH_3$); in the former the element is the basylous radical of acids (*e. g.*, HCl, $H_2SO_4$). In compound radicals, *e. g.*, $C_2H_3O_2$ or $NH_4$, the properties of hydrogen are no longer apparent; the chemical force resident in the atoms of such radicals seems to be mainly exerted in binding those atoms together.

Examples of Formulæ containing Univalent, Bivalent, and Trivalent Radicals.

The reader will find instructive practice in writing twenty or thirty imaginary formulæ of salts by placing in juxtaposition acidulous and basylous radicals, as in the following examples. Just as in a pair of scales a 2-lb. weight must be balanced by two 1-lb. weights, or a 4-lb. weight by two 2-lb. weights, or by one 3-lb. and one 1-lb. weight, so a bivalent radical unites with a bivalent radical or with two univalent radicals, a quadrivalent radical with two bivalent radicals, or with one trivalent and one univalent radical, and so on.

(R = any basylous Radical.) (*R* = any acidulous Radical.)

| *General formula.* | *Examples.* |
|---|---|
| $R'R'$ . . . . . | KI, NaCl, $NH_4C_2H_3O_2$, $AgNO_3$. |
| $R''R'_2$ . . . . | $CaCl_2$, $Zn2C_2H_3O_2$, $Pb2NO_3(BaNO_3C_2H_3O_2)$. |
| $R'''R'_3$ . . . . | $Bi3NO_3$, $AsH_3$, $SbCl_3$. |
| $R'_2R''$ . . } | $K_2CO_3$, $Na_2SO_4$, $H_2C_2H_4O_6$. |
| $R'R'R''$ . . } | $KHCO_3$, $NaHSO_4$, $KHC_4H_4O_6$. |
| $R'_3R'''$ . . } | $Am_3PO_4$, $K_3C_6H_5O_7$, $H_3AsO_3$. |
| $R'_2R'R'''$ . } | $Na_2HPO_4$, $Na_2HAsO_4$. |
| $R''R''$ . . . . | $CaCO_3$, MgO, $CuSO_4$, HgO, $FeSO_4$. |
| $R''R'''_2$ . . . . | $Ca_32PO_4$, $Ca_32C_6H_5O_7$. |
| $R''R'R'''$ . . . | $MgAmPO_4$, $CuHAsO_3$. |
| $R'''R''R'$ . . . | $BiONO_2$. |
| $R'''_2R''_2R''$ . . . | $Bi_2O_2CO_3$. |
| $R'''_2R''_3$ . . . . | $As_2O_3$, $Sb_2O_3$, $Fe_2O_3$, $Fe_23SO_4$. |
| $R'''R'''$ . . . . | $BiC_6H_5O_7$. |
| $R'''_2R'_6$ . . . . | $Fe_2Cl_6$, $Fe_26NO_3$, $Fe_26C_2H_3O_2$. |

Quadrivalent Radicals or *Tetrads*, Quinquivalent Radicals or *Pentads*, and Sexivalent Radicals or *Hexads*, are known.

---

## EXERCISE.

183. Write an exposition of the doctrine of Quantivalence within the limits of a sheet of note paper.

---

### DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE METALS, BARIUM, CALCIUM, MAGNESIUM.

Add yellow chromate of potassium to a portion of the solution to be examined; a precipitate indicates barium.

If no barium is present, add chloride and carbonate of ammonium, and boil; a precipitate indicates calcium.

If barium and calcium are proved to be absent, add chloride of ammonium, ammonia, and then either phosphate of sodium or arseniate of ammonium; a white granular precipitate indicates magnesium.

Ammonia is here added to yield the necessary elements to ammonio-magnesian phosphate or ammonio-magnesian arseniate, both of which are highly characteristic precipitates; and chloride of ammonium is added to prevent a mere partial precipitate of the magnesium by the ammonia.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF ONE, TWO, OR ALL THREE OF THE METALS, BARIUM, CALCIUM, MAGNESIUM.

Add chromate of potassium to the solution; barium, if present, is precipitated. Filter, if necessary, and add to the *filtrate* (that is, the liquid which has run through the filter) chloride, hydrate, and carbonate of ammonium, and boil; calcium, if present, is precipitated. Filter, if requisite, and add phosphate of sodium; magnesium, if present, is precipitated.

*Note.*—Red chromate of potassium must not be used in these operations, or a portion of the barium will remain in the liquid and be thrown down with, or in place of, the carbonate of calcium (*vide* p. 103). The yellow chromate must not contain carbonate of potassium, or calcium will be precipitated with, or in place of, barium. The absence of carbonate is proved by the non-occurrence of effervescence on the solution of hydrochloric acid to a little of the solution of the chromate, previously made hot in a test-tube. If the yellow chromate has been prepared by adding excess of ammonia to solution of red chromate of potassium, its addition to the liquid to be analyzed must be preceded by that of solution of chloride of ammonium, the precipitation of a portion of the magnesium (by the free ammonia in the yellow chromate) is thus prevented, for chloride of ammonium solution is a good solvent of hydrate (and carbonate) of magnesium, as already stated on page 120.

*Note* 1.—The analysis of solutions containing the foregoing metals is commenced by the addition of chloride of ammonium (AmCl) and ammonia (AmHO), simply as a precautionary measure, the former compound preventing partial precipitation of magnesium, the latter neutralizing acids. The carbonate of ammonium ($Am_2CO_3$) is the important group reagent—the precipitant of barium and calcium.

*Note* 2.—In the following, and in subsequent charts of analytical processes, the leading precipitants will be found to be ammonium salts. These, being volatile, can be got rid of towards the end of the operations, and thus the detection of potassium and sodium be in no way prevented—an advantage which could not be had if such salts as chromate of potassium or phosphate of sodium were the group-precipitants employed.

*Note* 3.—Acetic, and not hydrochloric or nitric acid is used in dissolving the barium and calcium carbonates, because chromate of barium, on the precipitation of which the detection of barium depends, is soluble in the stronger acids, and therefore could not be thrown down in their presence.

TABLE OF SHORT DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS CONTAINING **ANY OR ALL** OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.

To the solution add AmCl, AmHO, $Am_2CO_3$; boil and filter.

<table>
<tr><td colspan="2">Precipitate<br>Ba Ca.<br>Wash, dissolve in $HC_2H_3O_2$, add $K_2CrO_4$, and filter.</td><td colspan="2">Filtrate<br>Mg Am Na K.<br>Add $Am_2HPO_4$, shake. filter.</td></tr>
<tr><td>Precipitate<br>Ba.*</td><td>Filtrate<br>Ca.<br>Test by $Am_2C_2O_4$.</td><td>Precipitate<br>Mg.</td><td>Filtrate<br>Am Na K.<br>Evap. to dryness, ignite, dissolve residue in water.<br>Test for K by $PtCl_4$.<br>Test for Na by flame.<br>Test orig. sol. for Am.</td></tr>
</table>

*Note on Classification.*—The compounds of barium, calcium, and magnesium, like those of the alkali metals, have many analogies; the carbonate, phosphate, and arseniate of each is insoluble in water, which sufficiently distinguishes them from the members of the class first studied. They possess, however, well marked differences, so that their separation from each other is easy. The solubility of their hydrates in water marks their connection with the alkali metals; the slightness of that solubility, diminishing as we advance further and further from the alkalies, baryta being most and magnesia least soluble in water, points to their connection with the next class of metals, the hydrates of which are insoluble in water. These considerations must not, however, be over-valued. Though the solubility of their hydrates places barium nearest and magnesium furthest

* It is perhaps scarcely necessary to state that this precipitate is chromate of barium ($BaCrO_4$), as any reader who has carefully gone through the "foregoing analytical reactions" will know. The occurrence of chromate of barium at this particular place, however, and under the circumstances described, is abundant evidence of the presence of barium (in some form or other) in the liquid analyzed—which was a part of the problem to be solved by the operator. Similar remarks apply, of course, to the Ca, which is finally precipitated as oxalate ($CaC_2O_4$), to Mg, which is thrown out as ammonio-phosphate ($MgAmPO_4$), to Am, Na, and K, and to the elements similarly alluded to in the other subsequent tables for "short" directions for analysis.

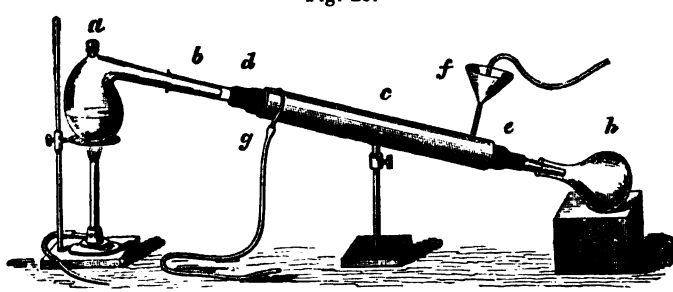
from the alkali metals, the solubility of their sulphates gives them the opposite order, magnesium-sulphate being most soluble, calcium-sulphate next, strontium-sulphate third (strontium is a rarer element, which will be mentioned subsequently), and barium-sulphate insoluble in water. These elements are sometimes spoken of as the metals of the alkaline earths.

*Note.*—In connection with the bivalence of the metals Barium, Calcium, and Magnesium, it is interesting to note that just as bivalent acidulous radicals give salts containing two atoms of univalent basylous radicals ($K_2SO_4$, $NaHSO_4$, $H_2CO_3$, $KNaC_4H_4O_6$), so bivalent basylous radicals yield salts containing two atoms of univalent acidulous radicals, as seen in acetonitrate of barium, $BaC_2H_3O_2NO_3$, a salt which is a definite compound, and not a mere mixture of acetate with nitrate of barium. A very large number of such salts is known.

## *Distillation.*

The water with which, in analysis, solution of a salt or dilution of a liquid is effected should be pure. Well or river-water (*Aqua*, U. S. P.) is unfit for the purpose, because containing alkaline and earthy salts (about 20 to 60 grains per gallon), derived from the soil through which the water percolates, and rain-water is not unfrequently contaminated with the dust and débris which fall on the

Fig. 28.




roofs whence it is usually collected. Such water is purified by *distillation*, an operation in which the water is by ebullition converted into steam, and the steam condensed again to water in a separate vessel, the fixed earthy and other salts remaining in the vessel in which the water is boiled. On the large scale, ebullition is effected in metal boilers having a hood or head in which is a lateral opening through which passes the steam; on the small scale, either a common glass flask is employed, into the neck of which, by a cork, is inserted a glass tube bent to an acute angle, or a *retort* is used (*a*, Fig. 28) a sort of long-necked Florence flask, dexterously bent near the body by the glass-worker to an appropriate angle (hence the name *retort*, from *retorqueo*, to bend beck). *Condensation* is effected by sur-

rounding the lateral steam-tube with cold water. In large stills the steam-tube, or *condensing-worm*, is usually a metal (tin) pipe, twisted into a spiral form for the sake of compactness, and so fixed in a tub that a few inches of one end of the pipe may pass through and closely fit a hole bored near the bottom of the tub. Cold water is kept in contact with the exterior of the pipe, provision being made for a continuous supply to the bottom, while the lighter water heated by the condensing steam runs off from the top of the column. The condenser for a flask or retort may be a simple glass tube of any size, placed within a second much wider tube (a common long, narrow lamp-glass answers very well for experimental operations), the inner tube being connected at the extremities of the wider by bored corks; a stream of water passes into one end of the inclosed space (the end furthest from the retort), through a small glass tube inserted in the cork, and out at the other through a similar tube. The common (Liebig's) form of laboratory condenser is a glass tube three-fourths of an inch wide and a yard long (*b*, Fig. 28), surrounded by a shorter tin or zinc tube (*c*, Fig. 28) two inches in diameter, and having at each extremity a neck, through which the glass tube passes. The ends of the necks of the tin tube, and small portions of glass tube near them, are connected by means of a strip of sheet caoutchouc carefully bound round, or by short, wide India-rubber tubes (*d* and *e*, Fig. 28). An aperture near the lower part of the tin tube provides for the admission of a current of cold water, and a similar aperture near the top (*g*, Fig. 28) allows the escape of heated water. The inner tube may thus constantly be surrounded by cold water, and heated vapors passing through it be perfectly cooled and condensed in any receiver (*h*, Fig. 28).

In distilling several gallons of water for analytical or medicinal purposes (*Aqua Destillata*, B. P. and U. S. P.), the first two or three pints should be rejected, because likely to contain ammoniacal and other volatile impurities.

*Rectification* is the process of redistilling a distilled liquid. *Rectified spirit* is spirit of wine thus treated.

*Dry* or *destructive distillation* is distillation in which the condensed products are directly formed by the decomposing influence of the heat applied to the dry or non-volatile substances in the retort or still.

---

## EXERCISE.

184. Write from memory two or three paragraphs descriptive of *distillation*.

---

## RECAPITULATION.

The subject just alluded to (distillation) naturally excites wonder respecting the cause of the physical difference between solid, liquid, and gaseous water. (Common observation will

have suggested to the student that the force of heat has much to do with the difference, and if he will turn to the chapter on *latent heat* in any book on Physics he will find that, as already indicated (p. 84), when ice liquefies by heat a very large amount of heat must be used before the slightest rise of temperature occurs. Afterwards the addition of heat makes the water hotter and hotter until one other point is reached (the boiling-point), when here again a great amount of heat is absorbed without causing the slightest rise in temperature. Afterwards more heat makes the gaseous water hotter and hotter, until, like a bar of iron, the steam, under special conditions, is made red-hot or white-hot. Different bodies absorb different amounts of heat in changing their physical condition from solid to liquid or liquid to gas (or vapor). The amount is constant for any one body, hence definite comparative numbers may be used for expressing the latent heats of substances.

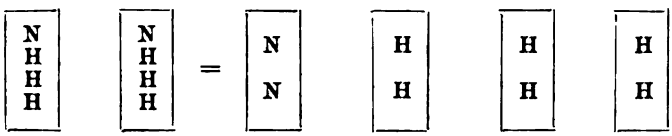
This absorption of heat at particular (liquefying and vaporizing) points, must not be confounded with an analogous physical action, namely, the absorption of heat which goes on when a body is rising in temperature. The amount of this absorption differs with different substances. That is to say, if equal weights of several substances, all at the same temperature, be all heated to a stated higher temperature, very different amounts of fuel will be required. The particular or specific amount in each case is always the same, hence the *specific heats* of substances may be expressed by numbers. See the chapter on "Specific Heat" in any Manual of Physics.

But after reading what has been stated respecting the constitution of matter (pp. 42 to 45), the chemical student will, in connection with the subject of distillation, be led, once more, to think over the subject of the molecular constitution of solid, liquid, and gaseous water, and of the molecular condition of bodies generally. As previously stated, little can be told him respecting the molecular condition of solids and liquids, for temperature and pressure affect them unequally, whence we conclude that, though the relation to each other of the molecules of any one substance is constant, this relation is different in different bodies. Different *gases*, however, are not differently affected but similarly affected by temperature and pressure, whence we conclude that their molecular constitution—the relations of their molecules to one another—is similar.

Another gas, ammonia, has been brought before the reader since the molecular constitution of gases was considered.

A small quantity of ammonia gas inclosed in the upper part of a roughly graduated test-tube over *mercury* (*water* would

dissolve it) and exposed to the continuous action of the electric spark, by means of wires of platinum fused in the sides of the tube, is decomposed into its elements nitrogen and hydrogen, the bulk of gas operated on being exactly doubled. This expansion is not due to the gaseous molecules receding from each other, but to every two molecules becoming four similar-sized molecules:—




Here each space (rectangular chiefly for convenience in printing) represents a molecule, and each letter one atom. Each space, if regarded as the side of a double cube, may also, for the moment, represent two volumes—such two volumes yielding, in the decomposition, one volume of nitrogen and three volumes of hydrogen, or the four such volumes of ammonia shown in the diagram yielding two volumes of nitrogen and six volumes of hydrogen.

Remembering that a symbol (of a gas) represents one volume, and that *a formula (of a gas) always represents two volumes*, the pupil will now see how full of meaning is such an equation as the following, including, as it does, names of the elements, number of the atoms, nature of the molecules, number of the molecules, weights of atoms of the molecules, and therefore weights of bulks of the bodies, and extent of expansion in the disunion of the elements, and therefore their extent of contraction in the act of union:—

$$2NH_3 = N_2 + 3H_2.$$

---

AT THIS STAGE THE LEARNER IS AGAIN RECOMMENDED TO READ THE PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY, AND TO RETURN TO THEM FROM TIME TO TIME UNTIL THEY ARE THOROUGHLY COMPREHENDED.

---

## ZINC, ALUMINIUM, IRON.

These three elements are classed together for analytical convenience rather than for more general analogies.

## ZINC.

Symbol Zn. Atomic weight 65.

*Source.*—Zinc is tolerably abundant in nature as sulphide (ZnS) or *blende*, and carbonate ($ZnCO_3$), or *calamine* (from *calamus*, a reed, in allusion to the appearance of the mineral). The ores are roasted to expel sulphur, carbonic acid gas, and some impurities, and the resulting oxide distilled with charcoal, when the metal vaporizes and readily condenses. Zinc is a brittle metal, but at a temperature somewhat below 300° F. is malleable, and may be rolled into thin sheets. Above 400° it is again brittle, and may then be pulverized. At 773° F. it melts, and at a bright red heat is volatile. Zinc in exceptionally fine powder ignites spontaneously, especially if damp or if stored in a warm place.

*Uses.*—Its use as a metal is familiar; alloyed with nickel and copper it yields german silver, with twice its weight of copper forms common brass, and as a coating on iron (the so-called *galvanized* iron) greatly retards the formation of rust. Most of the salts of zinc are prepared directly or indirectly from the metal (*Zincum*, B. P. and U. S. P.).

*Quantivalence.*—The atom of zinc is bivalent, Zn″.

*Molecular Weight.*—Some remarks on this point will be made under Mercury.

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Synthetical Reactions.*

#### Sulphate of Zinc.

*First Synthetical Reaction.*—Heat zinc (4 pts.) with water (20 pts.) and sulphuric acid (3 fl. pts.) in a test-tube (or larger vessel) until gas ceases to be evolved; solution of sulphate of zinc ($ZnSO_4$) results. Filter, to separate the particles of lead, carbon, etc., commonly contained in sheet zinc, and concentrate the solution in an evaporating dish; on cooling, colorless, transparent, prismatic crystals of Sulphate of Zinc ($ZnSO_4,7H_2O$) are deposited (*Zinci Sulphas*, B. P. and U. S. P.).

$$\underset{\text{Zinc.}}{Zn_2} + \underset{\text{Sulphuric acid.}}{2H_2SO_4} + \underset{\text{Water.}}{xH_2O} = \underset{\text{Sulphate of zinc.}}{2ZnSO_4} + \underset{\text{Water.}}{xH_2O} + \underset{\text{Hydrogen.}}{2H_2}$$

Zinc does not displace hydrogen from the sulphuric acid alone, nor from the water alone, yet the mixture affords hydrogen. The probable explanation is that as sulphuric acid combines with several different quantities of water to form definite hydrous compounds ($H_2SO_4,H_2O$; $H_2SO_4,2H_2O$; etc.), it is one of these that is de-

composed with elimination of hydrogen. At present we can only say that an unknown ($x$) amount of water is required in the reaction.

*Note.*—This reaction affords hydrogen and sulphate of zinc; it also develops electricity. Of several methods of evolving hydrogen, it is the most convenient; of the two or three means of preparing sulphate of zinc, it is the most commonly employed; and of the many reactions which may be utilized in the development of dynamic electricity, it is at present the cheapest and most manageable. The apparatus in which the reaction is effected differs according to the requirements of the operator; if the sulphate of zinc alone is wanted, an open dish is all that is necessary, the action being, perhaps, accelerated by heat; if hydrogen, a closed vessel and delivery-tube; if electricity, square vessels called cells, and certain complementary materials, forming altogether what is termed a battery. In each operation for one product the other two are commonly wasted. It would not be difficult for the operator, as a matter of amusement, to construct an apparatus from which all three products should be collected.

*Purification.*—Impure sulphate of zinc may be purified in the same manner as impure chloride (see next reaction).

Sulphate of zinc is isomorphous with sulphate of magnesium, and like that salt, loses six-sevenths of its water of crystallization at 212° F.

## Chloride of Zinc.

*Second Synthetical Reaction.*—Dissolve zinc in hydrochloric acid mixed with half its bulk of water; the resulting solution contains chloride of zinc. Evaporate the liquid till no more steam escapes; Chloride of Zinc ($ZnCl_2$) in a state of fusion remains, and on cooling, is obtained as an opaque white solid (*Zinci Chloridum*, B. P. and U. S. P.).

It is soluble in water, alcohol, or ether.

$$\underset{\text{Zinc.}}{Zn_2} + \underset{\text{Hydrochloric acid.}}{4HCl} = \underset{\text{Chloride of zinc.}}{2ZnCl_2} + \underset{\text{Hydrogen.}}{2H_2}$$

This reaction is analogous to that previously prescribed. The Burnett deodorizing or disinfecting liquid is solution of chloride of zinc.

*Purification of Chloride or Sulphate of Zinc.*—Zinc sometimes contains traces of iron or lead; and these, like zinc, are dissolved by most acids, with formation of soluble salts; they may be recognized in the liquids by applying the tests described hereafter, to a little of the solution in a test-tube. Should either be present in the above solution, a little chlorine water is added to the liquid till the odor of chlorine is permanent, and then the whole well shaken with some hydrate of zinc or the common official "carbonate" of zinc (really hydrato-carbonate—see p. 132). In this way iron is precipitated as ferric hydrate, and lead as peroxide:—

$$\underset{\text{Ferrous chloride.}}{*2FeCl_2} + \underset{\text{Chlorine.}}{Cl_2} = \underset{\text{Ferric chloride.}}{*Fe_2Cl_6}$$

$$\underset{\text{Ferric chloride.}}{Fe_2Cl_6} + \underset{\text{Carbonate of zinc.}}{3ZnCO_3} + \underset{\text{Water.}}{3H_2O} = \underset{\text{Ferric hydrate.}}{Fe_2 6HO} + \underset{\text{Chloride of zinc.}}{3ZnCl_2} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

$$\underset{\text{Chloride of lead.}}{PbCl_2} + \underset{\text{Chlorine.}}{Cl_2} + \underset{\text{Carbonate of zinc.}}{2ZnCO_3} = \underset{\text{Peroxide of lead.}}{PbO_2} + \underset{\text{Chloride of zinc.}}{2ZnCl_2} + \underset{\text{Carbonic acid gas.}}{2CO_2}$$

In the British Pharmacopœia the presence of impurities in the zinc is assumed, and the process of purification just described incorporated with the process of preparation of *Zinci Chloridum*, *Liquor Zinci Chloridi*, and *Zinci Sulphas*. In the purification of the sulphate of zinc, the action of chlorine on any ferrous sulphate will result in the formation of ferric sulphate as well as ferric chloride:—

$$6FeSO_4 + Cl_6 = 2(Fe_2 3SO_4) + Fe_2Cl_6;$$

carbonate of zinc will then give chloride as well as sulphate of zinc, and thus the whole quantity of sulphate of zinc be slightly contaminated by chloride. On evaporating and crystallizing, however, the chloride of zinc will be retained in the mother-liquor. This process admits of *general* application.

For *Liquor Zinci Chloridi*, B. P., 1 pound of zinc is placed in a mixture of 44 fluidounces of hydrochloric acid and 20 of water, the mixture ultimately warmed until no more gas escapes, filtered into a bottle, chlorine water added until the liquid after shaking smells fairly of chlorine, about half an ounce or somewhat more of carbonate of zinc shaken up with the solution until a brown precipitate (of ferric hydrate or peroxide of lead, or both) appears, the whole filtered, and the filtrate evaporated to 40 fluidounces. One fluidounce contains 366 grains of chloride of zinc. If there is reason to believe that neither iron nor lead is present in the zinc, the treatment with chlorine water and carbonate of zinc may be omitted. The *Liquor Zinci Chloridi*, U. S. P., is prepared by a somewhat similar process, nitric acid, however, is used instead of chlorine water; the solution contains 375 grains of chloride of zinc in one fluidounce.

## Carbonate of Zinc.

*Third Synthetical Reaction.*—To solution of any given quantity of sulphate of zinc in twice its weight of water (in a test-tube, evaporating basin, or other large or small vessel), add about an equal quantity of carbonate of sodium, also dissolved in twice its weight of water, and boil; the resulting white precipitate is so-called Carbonate of Zinc (*Zinci Carbonas*, B. P., *Zinci Carbonas Præcipitata*, U. S. P.), a mixture of carbonate ($ZnCO_3$) and hydrate (Zn2HO), in the pro-

* It will be noticed that the iron is represented, in these equations, as exerting both bivalent and trivalent activity; this will be alluded to when iron comes under consideration.

portion of one molecule of the former and two of the latter, together with a molecule of water ($H_2O$). It may be washed, drained, and dried in the usual manner. It is used in the arts under the name of *zinc-white*, and frequently in medicine in the form of ointment (*Ceratum Zinci Carbonatis*, U. S. P.).

$$3ZnSO_4 + 2H_2O + 3Na_2CO_3 = ZnCO_3,2ZnH_2O_2 + 2CO_2 + 3Na_2SO_4$$

Sulphate of zinc. — Water. — Carbonate of sodium. — Official carbonate of zinc. — Carbonic acid gas. — Sulphate of sodium.

## Acetate of Zinc.

*Fourth Synthetical Reaction.*—Collect in a filter the precipitate obtained in the last reaction, wash with distilled water, and dissolve a portion in strong acetic acid; the resulting solution contains acetate of zinc ($Zn2C_2H_3O_2$), and, on evaporating, and setting aside for a day, yields lamellar pearly crystals ($Zn2C_2H_3O_2,2H_2O$). This is the process for *Zinci Acetas*, B. P.

$$ZnCO_3,2ZnH_2O_2 + 6HC_2H_3O_2 = 3(Zn2C_2H_3O_2) + 5H_2O + CO_2$$

Official carbonate of zinc. — Acetic acid. — Acetate of zinc. — Water. — Carbonic acid gas.

The U. S. P. process consists in digesting oxide of zinc in acetic acid, heating the mixture to boiling point, filtering while hot, and setting aside the clear solution to crystallize.

## Oxide of Zinc.

*Fifth Synthetical Reaction.*—Dry the remainder of the precipitated carbonate (by placing the open filter on a plate over a dish of water kept boiling), and then heat it in a small crucible till it ceases to effervesce on the addition of water and acid to trial samples taken out of tbe crucible from time to time; the product is Oxide of Zinc (*Zinci Oxidum*, B. P. and U. S. P.), much used in the form of ointment (*Unguentum Zinci*, B. P., and *Unguentum Zinci Oxidi*, U. S. P.).

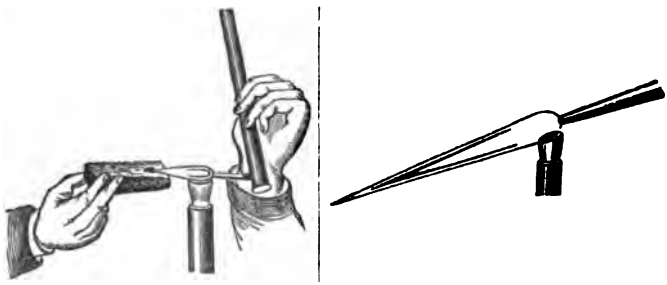
$$ZnCO_3,2ZnH_2O_2 = 3ZnO + 2H_2O + CO_2$$

Official carbonate of zinc. — Oxide of zinc. — Water. — Carbonic acid gas.

*Note.*—This oxide is yellow while hot, and of a very pale yellow or slight buff tint when cold, not actually white like the oxides prepared by the combustion of zinc in air. The latter variety occurs in

commerce under the name of Hubbuck's oxide of zinc. Its preparation can only be practically accomplished on the large scale, but the chief features of the action may be observed by heating a piece of zinc on charcoal in the blowpipe flame (Fig. 29) till it burns; flocks escape, float about in the air, and slowly fall. These are the old *Flores Zinci*, *Lana Philosophica*, or *Nihilum Album*.

Fig. 29. Fig. 30.



The Blowpipe.

A clear blowpipe-flame consists more or less of two portions (see Fig. 30), an inner cone at the apex of which are hot gases greedy of oxygen, and an outer cone at the apex of which is excess of hot oxygen. At the latter point oxidizable metals, etc., are readily oxidized, as in the foregoing experiment, and that part of the flame is therefore termed *the oxidizing* flame; in the inner flame oxides and other compounds (a grain of acetate of lead may be employed for illustration) are reduced to the metallic state, hence that part is termed *the reducing flame*. A blowpipe flame is much altered in character by slight variations in the position of the nozzle of the blowpipe, by the form of the nozzle, by the force with which air is expelled from the blowpipe, and by the character of the jet of gas.

## Valerianate of Zinc.

*Sixth Synthetical Reaction.*—Valerianate of Zinc ($Zn2C_5H_9O_2$) (*Zinci Valerianas*, B. P.) is prepared by mixing strong solutions of sulphate of zinc and valerianate of sodium, cooling, separating the white pearly crystalline matter, evaporating at 200° F. to a low bulk, cooling, again separating the lamellar crystals, washing the whole product with a small quantity of cold distilled water, draining and drying by exposure to air at

ordinary temperatures. Valerianate of zinc is soluble in ether, alcohol, or hot water.

$$\underset{\text{Sulphate of zinc.}}{ZnSO_4} + \underset{\text{Valerianate of sodium.}}{2NaC_5H_9O_2} = \underset{\text{Sulphate of sodium.}}{Na_2SO_4} + \underset{\text{Valerianate of zinc.}}{Zn2C_5H_9O_2}$$

*Note.*—The compounds of zinc described in the above six reactions are the only ones mentioned in the British Pharmacopœia; the processes are also those of that work. *Sulphide and Hydrate of Zinc* are mentioned in the following analytical paragraphs. The formula of *Sulphite of Zinc* is $ZnSO_3,3H_2O$.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—To solution of a zinc salt (sulphate, for example) in a test-tube, add solution of sulphydrate of ammonium ($NH_4HS$); white sulphide of zinc (ZnS) is precipitated, insoluble in acetic, but soluble in the stronger acids.

*Note.*—This is the only white sulphide that will be met with. Its formation, on the addition of the sulphydrate of ammonium, is therefore highly characteristic of zinc. If the zinc salt contains iron or lead as impurities, the precipitate will have a dark appearance, the sulphides of those metals being black. Hydrate of aluminium, which is also white and precipitated by sulphydrate of ammonium, is the only substance sulphide of zinc is likely to be mistaken for, and *vice versâ*; but, as will be seen immediately, there are good means of distinguishing these from each other.

*Second Analytical Reaction.*—To solution of a zinc salt add solution of ammonia; white hydrate of zinc (Zn2HO) is precipitated. Add excess of ammonia; the precipitate is redissolved.

This reaction at once distinguishes a zinc salt from an aluminium salt, hydrate of aluminium being insoluble in ammonia.

*Other Analytical Reactions.*—The fixed alkali-hydrates afford a similar reaction to that just mentioned, the hydrate of zinc redissolving if the alkali is free from carbonate.——Carbonate of ammonium yields a white precipitate of carbonate and hydrate, soluble in excess.——The fixed alkaline carbonates give a similar precipitate, which is not redissolved if the mixed solution and precipitate be well boiled.——Ferrocyanide of potassium precipitates white ferrocyanide of zinc ($Zn_2FeCy_6$).

Sulphate of magnesium, which is isomorphous with and indistinguishable in appearance from sulphate of zinc, is not precipitated from its solutions either by ferrocyanide of potassium or sulphydrate of ammonium.

*Antidotes.*—There are no efficient chemical means of counteracting the poisonous effects of zinc. Large doses, fortunately, act as powerful emetics. If vomiting has not occurred, or apparently to an insufficient extent, solution of carbonate of sodium (common washing salt) immediately followed by white of egg and demulcents may be administered.

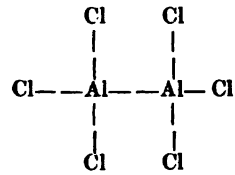
## QUESTIONS AND EXERCISES.

185. Give the sources and uses of metallic zinc.
186. Explain by a diagram what occurs when zinc is dissolved in diluted sulphuric acid.
187. How may solutions of Chloride and of Sulphate of Zinc be purified from salts of iron? Give equations descriptive of the reactions.
188. State the formula of Carbonate of Zinc, and illustrate by a diagram the reaction which takes place in its production.
189. Give an equation showing the formation of Acetate of Zinc.
190. In what respect does Oxide of Zinc, resulting from the ignition of the carbonate, differ from that produced during the combustion of the metal?
191. How is Valerianate of Zinc prepared?
192. What are the properties of Valerianate of Zinc?
193. Name the more important tests of Zinc.
194. How would you distinguish, chemically, between solutions of Sulphate of Zinc and Alum?
195. Describe the treatment in cases of poisoning by the salts of zinc.
196. Give reactions distinguishing Sulphate of Zinc from Sulphate of Magnesium.

## ALUMINIUM.

Symbol Al. Atomic weight 27.5.

*Note.*—In the formulæ of aluminium salts, it will be observed that to one atom of metal there are three atoms of other univalent radicals; hence, apparently, the atom of aluminium is trivalent, Al'''. But possibly it is quadrivalent; for one molecule of aluminium compounds includes two atoms of the metal, three-fourths only of whose power may be supposed to be exerted in retaining the other constituents of the molecule, the remaining fourth enabling the aluminium atoms themselves to keep together. This is graphically shown in the following formula of chloride of aluminium ($Al_2Cl_6$) from Frankland's "Lecture Notes for Chemical Students," which represents each



aluminium atom as a body having four arms or bonds, three of which are engaged in grasping the arms of univalent chlorine atoms, while the fourth grasps the corresponding arm of its brother aluminium atom. Such graphic formulæ, as they are called, are useful in facilitating the acquirement of hypotheses regarding the constitution of chemical substances, especially if the error be avoided of supposing that they are pictures either of the position or absolute power of atoms in a molecule, or indeed, the true representation of a molecule at all; for on this point man knows little or nothing.

*Source.*—Aluminium is very abundant in nature, chiefly as silicate, in clays, slate, marl, granite, basalt, and a large number of minerals. *Mica* consists chiefly of silicates of aluminium, iron, and potassium. *Rotten-stone* is an aluminium silicate containing a little organic matter. The sapphire and ruby are almost pure oxide of aluminium. The metal is obtained from the double chloride of aluminium and sodium, by the action of metallic sodium, the source of the chloride being the mineral *bauxite*—a more or less ferruginous hydrate of aluminium.

*Aluminium-bronze* is an alloy of ten parts of aluminium with ninety of copper.

*Alum* (*Alumen*, B. P. and U. S. P.), a double sulphate of aluminium and ammonium ($Al_2 3SO_4, Am_2SO_4, 24H_2O$), may be obtained from aluminous schist (from σχιστὸς, *schistos*, divided), a sort of pyritous slate or shale, by exposure to air; oxidation and chemical change produce sulphate of aluminium, sulphate of iron, and silica, from the silicate of aluminium and bisulphate of iron (iron pyrites) originally present in the shale. The sulphate of aluminium and sulphate of iron are dissolved out of the mass by water, and sulphate or chloride of ammonium added; on concentrating the liquid, alum crystallizes out, while the more soluble iron salt remains in the mother liquor.

Alum is also prepared by directly decomposing the silicate of aluminium in the calcined shale of the coal-measures by hot sulphuric acid, ammonia being added from time to time until a solution strong enough to crystallize is obtained. The liquid well agitated during cooling deposits alum, in minute crystals, termed *alum-flour*, which is afterwards recrystallized.

*Alums.*—There are several *alums*, iron or chromium taking the place of aluminium, and potassium or sodium that of ammonium, all crystallizing in an eight-sided form, the octahedron—a sort of double pyramid. These are apparently alike in chemical constitution, and their general formula (M = either metal) is $M'''_2 3SO_4, M'_2SO_4, 24H_2O$. The alum of the manufacturer commonly occurs in colorless, transparent, octahedral crystals, massed in lumps, which are roughly broken up for trade purposes, but still exhibit the faces of octahedra.

It contains sulphate of ammonium or sulphate of potassium according as one or other is the cheaper.

*Sulphate of Aluminium* ($Al_2 3SO_4, 9H_2O$), or *Alum Cake*, prepared from natural silicates in the manner just described, is a common article of trade, serving most of the manufacturing purposes for which alum was formerly employed. In the United States Pharmacopœia (*Aluminii Sulphas*) it is directed to be made by dissolving hydrate of aluminium in diluted sulphuric acid, with subsequent removal of water by evaporation.

$$Al_2 6HO + 3H_2SO_4 = Al_2 3SO_4 + 6H_2O.$$

The hydrate of aluminium is to be prepared by the addition of solution of alum to solution of carbonate of sodium, the precipitated hydrate being collected on a filter and well washed.

$$Al_2 3SO_4, Am_2SO_4 + 3Na_2CO_3 + 3H_2O = Al_2 6HO + Am_2SO_4 + 3Na_2SO_4 + 3CO_2.$$

*Preparation of Alum.*—Prepare alum by heating a small quantity of powdered pipe-clay (silicate of aluminium) with about twice its weight of sulphuric acid for some time, dissolving out the resulting sulphate of aluminium and excess of sulphuric acid by water, and adding ammonia to the clear filtered solution only until, after well stirring, the excess of acid is neutralized. (If too much ammonia be added, the hydrate of aluminium precipitated when the ammonia is first poured in will not be redissolved on well mixing the whole. Perhaps the readiest indication of neutrality in this and similar cases is the presence of a little precipitate after stirring and warming the mixture.) On evaporating the clear solution, crystals of alum are obtained.

The *Ammonio-ferric Alum* of American pharmacy (*Ferri et Ammonii Sulphas*, U. S. P.) is made by adding sulphate of ammonium to a hot solution of persulphate of iron, and setting the liquid aside to crystallize. It forms pale violet octahedral crystals expressed by the formula $Fe_2 3SO_4, (NH_4)_2SO_4, 24H_2O$. *Aluminii et Potassii Sulphas* (U. S. P.) is potassium alum, the sulphate of aluminium and potassium ($Al_2 3SO_4, K_2SO_4, 24H_2O$).

*Dried Alum* (*Alumen Exsiccatum*, B. P. and U. S. P.) is alum from which the water of crystallization has been expelled by heat, the temperature not exceeding 400° F. By calculation from the molecular weight of alum, it will be found that the salt contains between 47 and 48 per cent. of water. At temperatures above 400° alum is decomposed, sulphate of ammonium and sulphuric anhydride escaping, and pure oxide of aluminium ($Al_2O_3$) remaining. Dried alum rapidly reabsorbs water from the atmosphere. It is almost useless as a medicinal preparation.

*Roche alum*, or *Rock alum* (*roche*, French, rock), is the name of an impure native variety of alum containing iron. The article sold under this name is an artificial mixture of common alum with oxide of iron.

### Reactions having Analytical Interest.

*First Analytical Reaction.*—To a solution of an aluminium salt (alum, for example, which contains sulphate of aluminium) add sulphydrate of ammonium ($NH_4HS$); a gelatinous white precipitate of hydrate of aluminium falls:—

$$Al_2 3SO_4 + 6AmHS + 6H_2O = Al_2 6HO + 3Am_2SO_4 + 6H_2S.$$

*Second Analytical Reaction.*—To a solution of alum add ammonia, $NH_4HO$; hydrate of aluminium falls: add excess of ammonia; the precipitate is, practically, insoluble.

*Principle of Dyeing by help of Mordants.*—The precipitated hydrate of aluminium, or alumina, has great affinity for vegetable coloring-matters and also for the fibre of cloth. Once more perform the above experiment, but before adding the ammonia introduce some decoction of logwood, solution of cochineal, or other similar colored liquid, into the test-tube. Add now the ammonia, and set the tube aside for the alumina to fall; the latter takes down with it all the coloring principle. In dye-works the fabrics are passed through liquids holding the alumina but weakly in solution, and then through the coloring solutions; from the first bath the fibres abstract alumina, and from the second the alumina abstracts coloring-matter. Some other metallic hydrates, notably those of tin and iron, resemble alumina in this property; they are all termed *mordants* (from *mordens*, biting); the substances they form with coloring-matters have the name of *lakes*.

*Third Analytical Reaction.*—To the alum add solution of potash; again hydrate of aluminium falls. Add excess of potash, and agitate; the precipitate dissolves.

Hydrate of aluminium may be precipitated from this solution by neutralizing the potash with hydrochloric acid, and adding ammonia until, after shaking, the mixture has an ammoniacal smell, or by adding solution of chloride of ammonium to the potash liquid. But the former way is the better; for it is difficult to know when a sufficiency of the chloride of ammonium has been poured in, whereas reaction with blue and red litmus-paper at once enables the operator to know when excess of hydrochloric acid or ammonia has been added.

Alkaline carbonates, phesphates, arseniates, and salts of other acidulous radicals also decompose solutions of aluminium salts and produce insoluble compounds of that metal, with the several acidulous radicals (except the carbonic), but the resulting precipitates are of no special interest.

## QUESTIONS AND EXERCISES.

197. What is there remarkable about the quantivalence of aluminium?

198. Practically what is the quantivalence of the atom of aluminium?

199. Enumerate the chief natural compounds of aluminium.

200. Write down a formula which will represent either of the Alums.

201. Which *alum* is official, and commonly employed in the arts?

202. State the source, and explain the formation of alum.

203. What is the crystalline form of alum? Work a sum showing how much Dried Alum is theoretically producible from 100 pounds of alum. Ans. 52 lbs. 6 ozs.

204. Show by figures how ordinary ammonium alum is capable of yielding 11.356 per cent. of alumina.

205. Why are aluminium compounds used in dyeing?

206. How are salts of aluminium analytically distinguished from those of zinc?

---

## IRON.

Symbol Fe. Atomic weight 56.

*Sources.*—Compounds of iron are abundant in nature. ***Magnetic Iron Ore***, or *Loadstone* (*Lodestone* or *Leadstone*, from the Saxon *lædan*, to lead, in allusion to its use, or rather to the use of magnets made from it, in navigation), is the chief ore from which Swedish iron is made; it is a mixture of ferrous and ferric oxides ($FeO,Fe_2O_3$). Much of the Russian iron is made from *Specular Iron Ore* (from *speculum*, a mirror, in allusion to the lustrous nature of the crystals of this mineral). This and *Red Hæmatite* (from *αἷμα*, *haima*, blood, so named from the color of its streak), an ore raised in Lancashire, are composed of ferric oxide only ($Fe_2O_3$). *Brown Hæmatite*, an oxyhydrate, is the source of much of the French iron. *Spathic Iron Ore* (from *spatha*, a slice, in allusion to the lamellar structure of the ore) is a ferrous carbonate ($FeCO_3$). An impure ferrous carbonate forms the *Clay Ironstone*, whence most of the English iron is derived. The chief Scotch ore is also an impure carbonate, containing much bituminous matter; it is known as *Black Band*. *Iron Pyrites* (from *πῦρ*, *pur*, fire, in allusion to the production of sparks when sharply struck) ($FeS_2$) is a yellow lustrous mineral, of use only for its sulphur. Ferrous carbonate ($FeCO_3$), chloride ($FeCl_2,4H_2O$), and sulphate ($FeSO_4,7H_2O$) sometimes occur in springs, the water of which is hence termed *chalybeate* (*chalybs*, steel).

*Process.*—Iron is obtained from its ores by processes of roasting, and reduction of the resulting impure oxide with coal or charcoal in the presence of chalk, the latter uniting with the sand, clay, etc., to

form a fusible slag. The *cast iron* thus produced is converted into *wrought iron* (*Ferrum*, U. S. P.) by burning out the 4 or 5 per cent. of carbon, silicon, and other impurities present, by oxidation in a furnace, an operation which is termed *puddling*. *Steel* is iron containing from one to two per cent. of carbon, and is made either by burning out a portion of the carbon from cast iron or by strongly heating wrought iron in charcoal. The official variety of the metal (*Ferrum*, B. P.) is "wrought iron in the form of wire or nails free from oxide," the condition in which it is most easily employed for conversion into its compounds. In the form of a fine powder (see 17 Reac.) metallic iron is employed as a medicine.

*Properties.*—The specific gravity of pure iron is 7.844, of the best bar iron 7.7; its color is bluish-white or gray. Bar iron requires the highest heat of a wind-furnace for fusion, but below that temperature assumes a pasty consistence, and in that state two pieces may be joined or *welded* (Germ. *wellen*, to join) by the pressure of blows from a hammer. A little sand thrown on to the hot metal facilitates this operation by forming with the superficial oxide of iron a fusible slag, which is dispersed by the blows: the purely metallic surfaces are thus better enabled to come into thorough contact and enter into perfect union. Iron is highly ductile, and of all common metals possesses the greatest amount of tenacity. At a high temperature it burns in the air, forming oxide of iron. *Rust of iron* is chiefly red oxide of iron, with a little ferrous oxide and carbonate; it is produced by action of the moist carbonic acid of the air and subsequent oxidation. Steam passed over scrap iron heated to redness gives hydrogen gas and black oxide of iron. Iron exposed at a high temperature to oxidation by a limited amount of steam (Barff) or air (Bower) becomes coated with magnetic oxide of so closely coherent and adherent a nature that the metal is permanently protected from alteration by atmospheric and many other influences.

*Quantivalence.*—Iron combines with other elements and radicals in two proportions; those salts in which the atom of iron appears to possess inferior affinities (in which the other radicals are in the less amount) are termed *ferrous*, the higher being *ferric* salts. In the former the iron exerts bivalent ($Fe''$), in the latter trivalent activity ($Fe'''$ or $Fe_2^{VI}$).

The atom of iron is also sometimes considered to be sexivalent, on account of the analogy of its compounds with those of chromium, which is sexivalent, if the formula of its fluoride ($CrF_6$) be correct, and because the composition of *ferrate of potassium* ($K_2FeO_4$), a deep purple salt obtained on passing chlorine through a concentrated solution of potash in which fresh ferric hydrate is suspended, is best explained on the assumption of the sexivalence of its iron.

Why the quantivalence of the atom of iron should vary is not at present known.

*The Nomenclature of Iron Salts.*—For educational and descriptive purposes the two classes of iron compounds are very conveniently spoken of as *ferrous* and *ferric*, the syllable "*ferr*" common to all indicating their allied ferruginous character, the syllables *ous* and *ic* indicating the lower and higher class respectively—functions fulfilled

by these two syllables in other similar cases (sulphurous and sulphuric, mercurous and mercuric). Officially the iron salts are known by other names, thus, *Sulphate of Iron* (*Ferri Sulphas*), and *Phosphate of Iron* (*Ferri Phosphas*), names which are chemically inexplicit, for there are two sulphates, and two posphates, and the terms do not define which salt is intended. Consistency and uniformity would demand that the names Ferrous Sulphate, Ferrous Phosphate, or similar terms, should be employed. Practically, however, the old names cause no confusion, inasmuch as only one sulphate, phosphate, etc., are used in medicine; moreover, the higher salts usually have the prefix *per* attached (as persulphate, perchloride). These names are already well known, can be easily rendered in Latin, and then admit of simple abbreviations and adaptations such as are employed in prescriptions, advantages not possessed by the more rational terms. While, therefore, the comprehension of the chemistry of iron is rendered simple and intelligible by the use of the terms ferrous and ferric, the employment of older and less definite names may very well be continued in pharmacy as being practically more convenient.

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Synthetical Reactions.*

## FERROUS SALTS.

### Green Sulphate of Iron. Ferrous Sulphate.

*First Synthetical Reaction.*—Place iron (small tacks) in sulphuric acid diluted with eight times its bulk of water (in a test-tube, basin, or other vessel of any required size), accelerating the action by heat until effervescence ceases.

$$\underset{\text{Iron.}}{Fe_2} + \underset{\text{Sulphuric acid.}}{2H_2SO_4} = \underset{\text{Ferrous Sulphate.}}{2FeSO_4} + \underset{\text{Hydrogen.}}{2H_2}$$

The solution contains what is generally known as Sulphate of Iron, that is, Ferrous Sulphate, the lower of the two sulphates, and will yield crystals of that substance ($FeSO_4,7H_2O$) (*Ferri Sulphas*, B. P. and U. S. P.) on cooling or on further evaporation; or if the hot concentrated solutions be poured into alcohol, the mixture being well stirred, the sulphate is at once thrown down in minute crystals (*Ferri Sulphas Granulata*, B. P.). At a temperature of 400° F. ferrous sulphate loses six-sevenths of its water, and becomes the *Ferri Sulphas Exsiccata*, B. P. and U. S. P.

*Other Sources of Ferrous Sulphate.*—In the laboratory, ferrous sulphate is often obtained as a by-product in making sulphuretted hydrogen:—

$$FeS + H_2SO_4 = H_2S + FeSO_4.$$

In manufactories it occurs as a by-product in the decomposition of aluminous shale, as already noticed (p. 137).

Ten grains of granulated sulphate of iron dissolved in one ounce of water constitute "Solution of Sulphate of Iron," B. P. "The solution should be recently prepared."

*Notes.*—Ferrous sulphate is sometimes termed *green vitriol*. Vitriol (from *vitrum*, glass) was originally the name of any transparent crystalline substance, but afterwards restricted to the sulphates of zinc, iron, and copper, which were, and still are, occasionally known as white, green, and blue vitriol. *Copperas* (probably originally *Copper-rust*, a term applied to verdigris and other green incrustations of copper) is another name for this sulphate of iron, sometimes distinguished as *green copperas*, sulphate of copper being blue copperas. Solid sulphate of iron is a constituent of *Pilula Aloes et Ferri*, B. P.

Ferrous sulphate, when exposed to the air, gradually turns brown through absorption of oxygen, ferric oxysulphate ($Fe_2O2SO_4$) being formed. The latter is not completely dissolved by water, owing to the formation of a still lower insoluble oxysalt ($Fe_4O_5SO_4$) and soluble ferric sulphate, $5(Fe_2O2SO_4) = Fe_4O_5SO_4 + 3(Fe_2 3SO_4)$.

Iron heated with undiluted sulphuric acid gives sulphurous acid gas and ferrous sulphate:—

$$Fe_2 + 4H_2SO_4 = 2SO_2 + 2FeSO_4 + 4H_2O.$$

## Carbonate of Iron. Ferrous Carbonate.

*Second Synthetical Reaction.*—To solution of ferrous sulphate, boiling, in a test-tube, add a solution of carbonate of ammonium ($Am_2CO_3$, etc.) in recently boiled water; a white precipitate of ferrous carbonate ($FeCO_3$) is thrown down, rapidly becoming light green, bluish-green, and, after a long time, red, through absorption of oxygen, evolution of carbonic acid gas, and formation of ferric oxyhydrate.

| $FeSO_4$ | + | $Am_2CO_3$ | = | $FeCO_3$ | + | $Am_2SO_4$ |
|---|---|---|---|---|---|---|
| Ferrous sulphate. | | Carbonate of ammonium. | | Ferrous carbonate. | | Sulphate of ammonium. |

*Saccharated Carbonate of Iron.*—The above precipitate, rapidly washed with hot well-boiled distilled water, and the moist powder mixed with sugar, and quickly dried—in short, all possible precautions taken to avoid exposure to air—forms the saccharated carbonate of iron (*Ferri Carbonas Saccharata*, B. P.).

The official proportions are two ounces of the sulphate and one ounce and a quarter of the carbonate, each dissolved in half a gallon of hot water; the solutions are mixed and set aside in a deep well-

covered pan, the supernatant liquid poured off when the precipitate has subsided, the pan again filled up with boiling water, the liquid once more poured away, the precipitate transferred to a calico filter, drained, gently pressed, and while still somewhat moist rubbed in a mortar with one ounce of sugar, and finally dried over a water-bath.

Carbonate of iron, mixed with a fourth its weight of Confection of Roses (Honey and Sugar, U. S. P.), forms the *Pilula Ferri Carbonatis*, B. P. and U. S. P.

*Notes.*—The *Subcarbonate of Iron* (*Ferri Subcarbonas*, U. S. P.) is precipitated on mixing solutions of sulphate of iron and carbonate of sodium. On washing and drying it is converted into reddish-brown oxyhydrate of iron, water being absorbed and carbonic acid gas being eliminated. This oxyhydrate of iron is best made by precipitating solution of persulphate of iron by solution of soda, and washing and drying the product (see page 150).

Ferrous carbonate is said to be more easily dissolved in the stomach than any other iron preparation. It is so unstable and prone to oxidation, that it must be washed in water containing no dissolved air and mixed with the sugar (which protects it from oxidation) as quickly as possible. In making the official compound mixture of iron (*Mistura Ferri Composita*, B. P. and U. S. P.), "Griffith's mixture," the various ingredients, including the carbonate of potassium, should be placed in a bottle of the required size, space being left for the crystals or solution of ferrous sulphate, which should be added last, the bottle immediately filled up with rose-water, and securely corked; the minimum of oxidation is thus insured. *Pilulæ Ferri Compositæ*, U. S. P., is made from myrrh, carbonate of soda, sulphate of iron, and syrup; carbonate of iron is gradually formed.

$$\underset{\text{Ferrous sulphate.}}{FeSO_4} + \underset{\text{Carbonate of potassium.}}{K_2CO_3} = \underset{\text{Ferrous carbonate.}}{FeCO_3} + \underset{\text{Sulphate of potassium.}}{K_2SO_4}$$

## Arseniate of Iron. Ferrous Arseniate.

*Third Synthetical Reaction*, by which the lower arseniate of iron, ferrous arseniate (*Ferri Arsenias*, B. P.) ($Fe_32AsO_4$), partially oxidized, is formed. This will be noticed again under Arsenicum.

## Phosphate of Iron. Ferrous Phosphate.

*Fourth Synthetical Reaction.*—To solution of ferrous sulphate in a test-tube add a little solution of acetate of sodium, then solution of phosphate of sodium; the lower phosphate of iron, ferrous phosphate ($Fe_32PO_4$), is precipitated (*Ferri Phosphas*, B. P. and U. S. P.).

$$\underset{\text{Ferrous sulphate.}}{3FeSO_4} + \underset{\text{Phosphate of sodium.}}{2Na_2HPO_4} + \underset{\text{Acetate of sodium.}}{2NaC_2H_3O_2}$$
$$= \underset{\text{Ferrous phosphate.}}{Fe_2 2PO_4} + \underset{\text{Sulphate of sodium.}}{3Na_2SO_4} + \underset{\text{Acetic acid.}}{2HC_2H_3O_2}$$

Officially, solutions of 3 ounces of sulphate of iron in a quart of water and 2½ ounces of phosphate and 1 of acetate of sodium in another quart of water, are well mixed, filtered, the precipitate well washed, and, to prevent oxidation as much as possible, dried at a temperature not exceeding 120° F. These proportions will be found to accord with the molecular weights of the crystalline salts, multiplied as indicated in the foregoing equation. $3(FeSO_4,7H_2O)=834$; $2(Na_2HPO_4,12H_2O)=716$; $2(NaC_2H_3O_2,3H_2O)=272$.

The above reaction also occurs in making *Syrupus Ferri Phosphatis*, B. P. The precipitate should be well washed, or red ferric acetate may be developed after a time.

*The use of acetate of sodium* (not mentioned in the U. S. P. formula) is to insure the absence of free sulphuric acid in the solution —sulphate of sodium being formed together with acetic acid. Sulphuric acid is a powerful solvent of ferrous phosphate; acetic acid is only a weak solvent. It is impossible to prevent the separation of sulphuric acid, if only ferrous sulphate and phosphate of sodium be employed. The solvent action of the sulphuric acid for the ferrous phosphate is still better prevented by the use of excess of the phosphate of sodium (Price), sodium sulphate, and doubtless monosodium phosphate ($NaH_2PO_4,H_2O$) resulting. Ferrous phosphate is white, but soon oxidizes and becomes slate-blue.

## Sulphide of Iron. Ferrous Sulphide.

*Fifth Synthetical Reaction.*—In a gas- or spirit-flame strongly heat sulphur with about twice its weight of iron filings in a test-tube (or in an earthen crucible in a furnace); ferrous sulphide (FeS) is formed. When cold, add water to a small portion, and then a few drops of sulphuric acid; sulphuretted hydrogen gas ($H_2S$), known by its odor, is evolved.

$$FeS + H_2SO_4 = FeSO_4 + H_2S.$$

Sticks of sulphur pressed against a white-hot bar of cast iron give a pure form of ferrous sulphide. The liquid sulphide thus formed is allowed to drop into a vessel of water (Sulphide of Iron, B. P.; Ferri Sulphuretum, U. S. P.). Or melted sulphur may be poured into a crucible full of red-hot iron nails, when a quantity of fluid ferrous sulphide is at once formed and may be poured out on to a slab.

### Green Iodide of Iron. Ferrous Iodide.

*Sixth Synthetical Reaction.*—Place a piece of iodine, about the size of a pea, in a test-tube with a small quantity of water, and add a few iron filings, small nails, or iron wire. On gently warming, or merely shaking if longer time be allowed, the iodine disappears, and, on filtering, a clear light green solution of iodide of iron ($FeI_2$) is obtained.

*The official Ferri Iodidum* is formed by gently warming a mixture of 3 parts of iodine, 1½ of fine iron wire, and 12 of distilled water in an iron vessel. When combination is nearly complete (as shown by indications of a sea-green tint), boil for a short time until the whiteness of the froth proves that the iodine has entirely disappeared. The solution is then filtered and evaporated in a clean bright iron saucepan, ladle, or dish until a drop taken out on the end of an iron wire stirrer solidifies on cooling. The liquid is poured out on a clean smooth slab, broken up and preserved in a glass-stoppered bottle. Solid iodide of iron has a crystalline fracture, is "green with a tinge of brown; inodorous, deliquescent, and almost entirely soluble in water, forming a slightly green solution which gradually deposits a colored sediment and acquires a red color."

The solid iodide contains about 18 per cent. of water of crystallization, and a little oxide of iron. It is deliquescent and liable to absorb oxygen from the air with formation of insoluble ferric oxyiodide or hydrato-iodide. Iodide of iron thus spoiled may be purified by resolution in water, addition of a little more iodine and some iron, warming, filtering, and evaporating as before.

*Ferrous bromide* ($FeBr_2$), occasionally used in medicine, could be made, as might be expected, in the same way as the iodide.

## FERRIC SALTS.

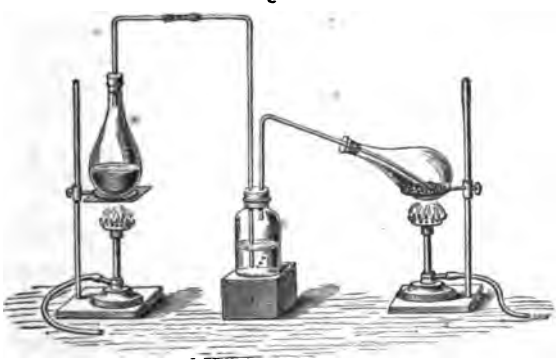
### Anhydrous Perchloride of Iron. Ferric Chloride.

*Seventh Synthetical Reaction.*—Pass chlorine (generated as usual, from black oxide of manganese and hydrochloric acid in a flask) through sulphuric acid contained in a small bottle, and thence by the ordinary narrow glass tubing quite to the bottom of a test-tube containing twenty or thirty small iron tacks (or a Florence flask containing 2 or 3 ounces of iron tacks), the latter kept hot by a gas-flame; the higher chloride of iron, ferric chloride, or the perchloride* of iron ($Fe_2Cl_6$) is formed, and condenses in the upper part of the tube or flask as a mass of small dark iridescent crystals. When a tolerably thick crust

* The prefixes *per* and *hyper* used here and elsewhere are from ὑπερ, *hyper*, over and above, and simply mean "the highest" of several. Thus perchloride, the highest *chloride*.

of the salt is formed, break off the part of the glass containing it, being careful that the remaining corroded tacks are excluded, and place it in ten or twenty times its weight of water; the resulting solution, poured off from any pieces of

Fig. 31.



Preparation of Anhydrous Ferric Chloride.

glass, is a pure neutral solution of hydrous ferric chloride, and will be serviceable in performing analytical reactions.

*Precaution.*—The above experiment must be conducted in the open air, or in a cupboard having a draught outwards.

*Anhydrous Ferrous Chloride.*—In breaking up the tube, small scales of a light buff color will be observed adhering to the nails; they are crystals of ferrous chloride ($FeCl_2$).

*Note.*—Solution of ferric chloride evolves some hydrochloric acid on boiling, while a darker colored solution of ferric oxychloride remains.

## Green Chloride of Iron. Hydrous Ferrous Chloride. Solution of Hydrous Ferric Chloride.

*Eighth Synthetical Reaction.*—Dissolve iron tacks, in a test-tube, in hydrochloric acid; hydrogen escapes, and the solution on cooling, or on evaporation and cooling, deposits *crystallized ferrous chloride* ($FeCl_2$), associated with four molecules of water ($4H_2O$) of crystallization ($FeCl_2,4H_2O$).

Through a portion of the solution of ferrous chloride pass chlorine gas; the ferrous chloride becomes *ferric chloride.*

The excess of chlorine dissolved by the liquid in this experiment may be removed by ebullition; but the ferric chloride is slightly de-

composed at the same time, for the reason just stated. The free chlorine may also be carried off by passing a current of air through the liquid for some time.

### Hydrous Ferric Chloride (another process).

*Ninth Synthetical Reaction.*—To another portion of the solution of ferrous chloride, in a test-tube, add a little more hydrochloric acid; heat the liquid, and continue to drop in nitric acid until the black color it first produces disappears; the resulting reddish-brown liquid is also solution of ferric chloride.

$$\underset{\text{Ferrous chloride.}}{6FeCl_2} + \underset{\text{Nitric acid.}}{2HNO_3} + \underset{\text{Hydrochloric acid.}}{6HCl} = \underset{\text{Ferric chloride.}}{3Fe_2Cl_6} + \underset{\text{Nitric oxide.}}{2NO} + \underset{\text{Water.}}{4H_2O}.$$

The black substance is a compound of nitric oxide gas (NO) with a portion of the ferrous salt; it is decomposed by heat.

This is the process for producing the *Liquor Ferri Perchloridi Fortior*, B. P., 2 ounces of iron, 12 fluidounces of hydrochloric acid, 9 fluidrachms of nitric acid, and 8 ounces of water being employed, and the product boiled down to 10 fluidounces. Practically it is impossible so to apportion the acids that a solution shall result containing neither excess of acid nor of metal, nor contain ferric nitrate. For most medicinal purposes, however, solution of perchloride of iron containing hydrochloric acid is said to be unobjectionable. The *Liquor Ferri Chloridi*, U. S. P., is a similar solution, but not quite so strong.

Diluted with 3 volumes of water this strong solution gives the *Liquor Ferri Perchloridi*, B. P.—or with 3 volumes of rectified spirit the *Tinctura Ferri Perchloridi*, B. P. The *Tinctura Ferri Chloridi*, U. S. P., is a similar solution.

*Note.*—The spirit in the tincture is unnecessary, useless, and deleterious; for it acts neither as a special solvent nor as a preservative, the offices usually performed by alcohol (*Tincturæ et Succi*, B. P. and U. S. P.); but, unless the liquid contain excess of acid, decomposes the ferric chloride and causes the formation of an insoluble oxychloride of iron. Even if the tincture be acid, it slowly loses color, ferrous chloride and chlorinated ethereal bodies being formed. A *Liquor*, of similar strength, is doubtless destined to displace the tincture altogether.

Solution of ferric chloride evaporated yields a mass of yellow crystals (*Ferri Chloridum*, U. S. P.) containing $Fe_2Cl_6,12H_2O$, or, rarely, red crystals having the formula $Fe_2Cl_6,5H_2O$. An old method of making solution of ferric chloride is to dissolve ferric oxide or hydrate in hydrochloric acid; but from the varying character of trade specimens of the ingredients, the liquid is more likely to contain excess or deficiency of iron than the proper proportion.

## Persulphate of Iron. Ferric Sulphate.

*Tenth Synthetical Reaction.*—Dissolve about three-quarters of an ounce of ferrous sulphate and a sixth of its weight of sulphuric acid in an ounce and a half of water in an evaporating dish, heat the mixture and drop in nitric acid until the black color it first produces disappears; the resulting liquid, when made of a certain prescribed strength, is the solution of ferric sulphate, or higher sulphate, "Solution of Persulphate of Iron" of the British Pharmacopœia, a heavy dark-red liquid, sp. gr. 1.441. *The Liquor Ferri Tersulphatis* U. S. P., is the same preparation, but slightly stronger (sp. gr. 1.320). *Liquor Ferri Subsulphatis*, U. S. P. (Monsel's solution), is a similar fluid, made with less acids, probably containing, therefore, ferric oxysulphate and ferric oxynitrate (sp. gr. 1.552).

$$\underset{\text{Ferrous sulphate.}}{6FeSO_4} + \underset{\text{Sulphuric acid.}}{3H_2SO_4} + \underset{\text{Nitric acid.}}{2HNO_3} = \underset{\text{Ferric sulphate.}}{3(Fe_2 3SO_4)} + \underset{\text{Nitric oxide.}}{2NO} + \underset{\text{Water.}}{4H_2O}$$

The black color, as in the previous reaction, is due to a compound of ferrous salt with nitric oxide ($2FeSO_4 + NO$).

*Note.*—In all the reactions in which iron passes from ferrous to ferric condition the element assumes different properties, the chief one being an alteration from bivalent to trivalent activity.

## Acetate of Iron. Ferric Acetate.

*Eleventh Synthetical Reaction.*—To a strong solution of ferric sulphate (from which free nitric acid has been removed by evaporating to dryness and redissolving in water) add an *alcoholic* solution of acetate of potassium ($KC_2H_3O_2$), and well shake the mixture; a crystalline precipitate of sulphate of potassium ($K_2SO_4$) falls, and ferric acetate ($Fe_2 6C_2H_3O_2$) remains in solution, forming, when filtered, and of definite strength, the *Tinctura Ferri Acetatis*, B. P. The preparation is unstable.

$$\underset{\text{Ferric sulphate.}}{Fe_2 3SO_4} + \underset{\text{Acetate of potassium.}}{6KC_2H_3O_2} = \underset{\text{Sulphate of potassium.}}{3K_2SO_4} + \underset{\text{Ferric acetate.}}{Fe_2 6C_2H_3O_2}$$

The official proportions are 2½ fluidounces of "Solution of Persulphate of Iron" with 8 fluidounces of rectified spirit, mixed with a solution of 2 ounces of acetate of potassium in 10 fluidounces of spirit, the whole well shaken frequently during an hour, filtered, and the precipitated sulphate of potassium washed by pouring on spirit until the filtrate measures 1 pint. A solution four times this strength, made from ferric hydrate and glacial acetic acid, is stable: it is diluted with spirit as wanted (J. Deane and T. Jeaffreson).

## Perhydrate of Iron. Ferric Hydrate.

*Twelfth Synthetical Reaction.*—Pour a portion of the solution of ferric sulphate into excess of solution of soda (ammonia, U. S. P.); moist ferric hydrate is precipitated (*Ferri Peroxidum Humidum*, B. P., *Ferri Oxidum Hydratum*, U. S. P.).

$$\underset{\text{Ferric sulphate.}}{Fe_2 3SO_4} + \underset{\text{Soda.}}{6NaHO} = \underset{\text{Ferric hydrate.}}{Fe_2 6HO} + \underset{\text{Sulphate of sodium.}}{3Na_2SO_4}.$$

Either of the other alkalies (potash or ammonia) will produce a similar reaction; but soda is cheapest and most convenient.

*Ferric hydrate is an antidote for arsenic* if administered directly the poison has been taken.

It converts the soluble arsenic ($As_2O_3$) into insoluble ferrous arseniate:—

$$2(Fe_2 6HO) + As_2O_3 = Fe_2 2AsO_4 + 5H_2O + Fe2HO.$$

Dried ferric hydrate (having become an oxyhydrate—$Fe_4O_4 4HO$) (*Ferri Peroxidum Hydratum*, B. P.) has less action on arsenic. Even the moist recently prepared hydrate ($Fe_2 6HO$) loses much of this power as soon as it has become converted into an oxyhydrate ($Fe_4O_3 6HO$), a change which occurs though the hydrate be kept under water (W. Procter, Jr). According to T. and H. Smith this decomposition occurs gradually, but in an increasing ratio; so that after four months the power of the moist mass is reduced to one-half and after five months to one-fourth. Now mere loss of water is not usually followed by any alteration of the essential chemical properties of a compound. It would seem, therefore, that ferric hydrate (two molecules) ($Fe_4 12HO$) probably suffers, on standing, actual decomposition into oxyhydrate ($Fe_4O_3 6HO$) and water ($3H_2O$), and does not merely lose water already existing in it as water. Ferric hydrate is also far more readily soluble in hydrochloric acid, tartaric acid, citric acid, and acid tartrate of potassium, than ferric oxyhydrate. Any formula exhibiting ferric hydrate ($Fe_2 6HO$) as a combination of ferric oxide and water ($Fe_2O_3, 3H_2O$) is, apparently, for these and other reasons, incorrect.

## Peroxyhydrate of Iron. Ferric Peroxyhydrate.

Collect the precipitate on a filter, wash, and dry on a plate over hot water; ferric oxyhydrate (*Ferri Peroxidum Hydratum*, B. P.) ($Fe_2O_2 2HO$) remains. When rubbed to powder it is fit for use in medicine.

$$Fe_2 6HO = Fe_2O_2 2HO + 2H_2O.$$

This oxyhydrate further decomposes when heated to low redness, ferric oxide ($Fe_2O_3$) remaining.

$$Fe_2O_2 2HO = Fe_2O_3 + H_2O.$$

## Peroxide of Iron. Ferric Oxide.

The six univalent atoms of the HO, the characteristic elements of all hydrates, are thus, by two successive steps, split up into water and oxygen. But between the hydrate and oxide there obviously may be another oxyhydrate, in which only 2HO is displaced by O″, and such a compound is well known; it is a variety of brown iron ore. The other oxyhydrate, $Fe_2O_2 2HO$, is also native (needle iron ore) as well as being the *Ferri Peroxidum Hydratum*, B. P.

| | |
|---|---|
| "Ferri Peroxidum Humidum" . . . . . | $Fe'''_2$ 6HO |
| A variety of brown iron ore . . . . . | $Fe'''_2 O''$ 4HO |
| "Ferri Peroxidum Hydratum" (needle ore) . | $Fe'''_2 O''_2$ 2HO |
| Ferric oxide . . . . . | $Fe'''_2 O''_3$ |

The moist ferric hydrate, as already stated, when kept for some months, even under water, loses the elements of water, and is converted into an oxyhydrate having the formula $Fe_4H_6O_9$ (limonite or brown hæmatite), which is either a compound of the above oxyhydrates $(Fe_2O4HO)+(Fe_2O_2 2HO)$, or is a definite intermediate oxydrate $(Fe_4O_5 6HO)$.

By ebullition with water for seven or eight hours, ferric hydrate is decomposed into water and an oxyhydrate having the formula $Fe_4H_2O_7$ (Saint Giles), which is either a mixture of the official oxyhydrate $(Fe_2O_2 2HO)$ with ferric oxide $(Fe_2O_3)$, or a definite intermediate body $(Fe_4O_5 2HO)$. The relation of these bodies to each other will be apparent from the following Table, in which, for convenience, the formulæ of ferric hydrate and oxide are doubled.

| | |
|---|---|
| Ferric hydrate (B. P.) (as stalactite) . . . | $Fe_4$ 12HO |
| Kilbride mineral (?) . . . . . | $Fe_4 O$ 10HO |
| Brown iron ore (Huttenrode and Raschau) . | $Fe_4 O_3$ 8HO |
| Old or frozen ferric hydrate (limonite) . . | $Fe_4 O_3$ 6HO |
| Ferric oxyhydrate (B. P.) (göthite) . . . | $Fe_4 O_4$ 4HO |
| Boiled ferric hydrate (turgite) . . . . . | $Fe_4 O_5$ 2HO |
| Ferric oxide (red hæmatite) . . . . . | $Fe_4 O_6$ |

A ferric oxycarbhydrate $(Fe_4OCO_3 8HO)$ has been obtained by Rother.

*The English official ferric oxyhydrate* $(Fe_2O_2 2HO)$, termed in the British Pharmacopœia *Hydrated Peroxide of Iron*, under the assumption that it is a compound of ferric oxide and water $(Fe_2O_3, H_2O)$, was formerly made by mixing solutions of ferrous sulphate and carbonate of sodium and exposing the resulting ferrous carbonate to the air until it was nearly all converted into ferric oxyhydrate; hence its old name, still sometimes seen on old bottles, of *Ferri Carbonas* and *Ferri Subcarbonas*.

## Ferric Oxide (another process).

*Thirteenth Synthetical Reaction.*—Roast a crystal or two of ferrous sulphate in a small crucible until fumes cease to be

evolved; the residue is a variety of ferric oxide ($Fe_2O_3$) or peroxide of iron, known in trade as *red oxide of iron, colcothar, crocus, rouge* (mineral), or *Venetian red.* It has sometimes been used in pharmacy in mistake for the official oxyhydrates (*vide* 12th Synthet. Reac.), from which it differs not only in composition but in the important respect of being almost insoluble in acids.

### The Scale Compounds of Iron.

*Fourteenth Synthetical Reaction.*—Repeat the twelfth reaction, introducing a little solution of citric or tartaric acid, or acid tartrate of potassium, before adding to the alkali (soda, potash, or ammonia), and notice that now no precipitation of ferric hydrate occurs. This experiment serves to illustrate, not the manufacture of a scale compound, but the chemistry of the manufacture. The effect is due to the formation of double compounds, termed Ammonio-Citrate, Potassio-Citrate, Ammonio-Tartrate, Potassio-Tartrate, and similar Sodium compounds of Iron, which remain in solution along with the secondary product—sulphate of the alkali metal. Such ferric compounds, made with certain prescribed proportions of recently prepared ferric hydrate (from which all alkaline sulphate has been washed), and the respective acids (tartaric or citric) or acid salts (acid tartrate of potassium), etc., and the solutions evaporated to a syrupy consistence, and spread on flat plates till dry, form the scale preparations known as *Ferri et Ammonii Citras*, U. S. P., *Ferri Citras*, U. S. P. (also *Liquor Ferri Citratis*, U. S. P.), *Ferri et Ammonii Tartras*, U. S. P., and *Ferri Potassio-tartras*, or rather *Ferrum Tartaratum*, B. P., *Ferri et Potassii Tartras*, U. S. P. A mixture of citrate of iron and ammonium with citrate of strychnia yields, on evaporation, *Ferri et Strychniæ Citras*, U. S. P. A mixture of ferric citrate with citrate of ammonium and citrate of quinine yields, by similar treatment, the well-known scales of *Ferri et Quiniæ Citras*, B. P. and U. S. P.

Specimens of these substances may be prepared by attending to the following details. It is essential, first, that the ferric hydrate be thoroughly washed, or an insoluble oxysulphate will be formed; second, that the ferric hydrate be rapidly washed, or an insoluble ferric oxyhydrate will be produced; thirdly, that the whole operation be conducted quickly, or reduction to green ferrous salt will occur; fourthly, that the solutions of the salts be not evaporated at a higher temperature than that stated, or decomposition will take place; and fifthly, that excess of ferric hydrate be employed.

In the pharmacopœial processes for the three scale compounds, the ferric hydrate is in each case freshly made from solution of ferric sulphate by precipitation with solution of ammonia.

$$\underset{\text{Ferric sulphate.}}{Fe_2 3SO_4} + \underset{\text{Hydrate of ammonium.}}{6AmHO} = \underset{\text{Ferric hydrate.}}{Fe_2 6HO} + \underset{\text{Suphate of ammonium.}}{3Am_2SO_4}$$

the solution of ferric sulphate being made of a definite strength from a known weight of ferrous sulphate. The reason for adopting this course is that ferric hydrate is unstable and cannot be weighed, because it cannot be dried without decomposing and becoming insoluble, as explained under the 12th reaction. This definite solution of ferric sulphate (*Liquor Ferri Persulphatis*, B. P., *Liquor Ferri Tersulphatis*, U. S. P.) is made by adding six fluidrachms of sulphuric acid to half a pint of water, warming, dissolving eight ounces of crystals of sulphate of iron in the liquid, pouring in nitric acid (six fluidrachms or rather more) slightly diluted until the mixture turns from a black to a reddish color, and ruddy nitrous vapors cease to be produced; the whole should measure eleven fluidounces, being diluted or further evaporated, as the case may be, to this bulk.

$$6FeSO_4 + 3H_2SO_4 + 2HNO_3 = 3(Fe_2 3SO_4) + 2NO + 4H_2O.$$

*Ferri et Ammoniæ Citras*, B. P., *Ferri et Ammonii Citras*, U. S. P.—Ferric hydrate is dissolved in solution of citric acid, ammonia added, and the whole evaporated to dryness.

To prepare the ferric hydrate, dilute eight or, rather, nine and a half fluidounces of the above solution of ferric sulphate with about a quart of water; pour this into two or three pints of water containing excess of solution of ammonia—about five fluidounces of "Strong Solution of Ammonia," or 15 ounces of "Solution of Ammonia." (If the opposite course were adopted, the alkaline liquid poured into the ferric solution, the precipitate would contain ferric oxysulphate, or hydrato-sulphate, which interferes with the brilliancy of the scales.) Thoroughly stir the mixture (it will smell strongly of ammonia, if enough of the latter has been added), allow the precipitate to subside, pour away the supernatant liquid, add more water, and repeat the washing until a little of the liquid tested for by-product (sulphate of ammonium) by solution of chloride or nitrate of barium ceases to give a white precipitate (sulphate of barium). Collect the ferric hydrate on a filter, drain, and place, while still moist, in a solution of four ounces of citric acid in eight of water, placed in an evaporating basin, over a water-bath, stir frequently, until the whole, or nearly the whole, of the hydrate has dissolved. To the solution, when cold, add nearly two fluidounces of strongest (or five and a half of weak) solution of ammonia, filter, evaporate over a water-bath to the consistence of syrup, spread thinly on panes of glass, and dry (at a temperature not exceeding 100° F.). The product scales off the glass in deep red transparent laminæ.

*Ferri et Quiniæ Citras*, B. P. and U. S. P.—Ferric hydrate and pure quinia are dissolved in solution of citric acid, ammonia added, and the whole evaporated to dryness. The product contains citrate of iron, citrate of quinia, and citrate of ammonium.

The ferric hydrate is obtained from four and a half fluidounces of the solution of ferric sulphate, with all the precautions described in the previous paragraph, a proportionate quantity of ammonia being employed.

While the ferric hydrate is being washed, prepare the quinia by dissolving one ounce of the ordinary sulphate of quinia in eight ounces of distilled water, acidified with sufficient sulphuric acid to dissolve the sulphate (about 12 fluidrachms of the official "diluted sulphuric acid"), and to the clear liquid add solution of ammonia, well mixing the product by stirring, until the whole of the quinia is precipitated (that is, until the mixture, after thorough agitation, smells of ammonia). Collect the precipitate on a filter, let it drain, and wash away adhering solution of sulphate of ammonium by passing through it about a pint and a half of distilled water.

(It will be observed that the principle involved in the preparation of quinia from its sulphate is identical with that which obtains in the precipitation of alumina, ferric hydrate, or hydrate of zinc, etc. Such a soluble sulphate—or, indeed, any similar soluble salt—has its acidulous constituent removed by the superior affinity of the basylous radical in ammonia, or other alkali, an insoluble precipitate and a new soluble sulphate being formed. The latter is washed away, leaving the former pure. In such manipulations, when economy has to be practised, soda is the alkali generally employed. Ammonia, however, has the advantage of showing the moment when its work of removing an acidulous radical is completed; for the salts which ammonium forms with such acidulous radicals as $SO_4$, Cl, $NO_3$, and $C_2H_3O_2$ are inodorous, while it itself has a powerful odor; so long, therefore, as the salt to be decomposed is not wholly attacked, the addition of ammonia does not give an ammoniacal odor to the mixture, the ammonia, as such, being, in fact, destroyed; but when the work is accomplished, the quantity of ammonia last added remains as ammonia, and communicates its natural smell to the liquid.)

The ferric hydrate and quinia being now washed and drained, dissolve the former, and afterwards the latter, in a solution of three ounces of citric acid in five of distilled water, the acid liquid being warmed over a water-bath, and portions of the precipitates stirred in as fast as solution is effected. "Let the solution cool, then add in small quantities at a time twelve fluidrachms of solution of ammonia diluted with two fluidounces of distilled water, stirring the solution briskly, and allowing the quinia which separates with each addition of ammonia to dissolve (in the acid) before the next addition is made. (Excess of ammonia must be avoided or the quinia will be precipitated.) Filter the solution, evaporate to the consistence of a thin syrup, and then dry in thin layers on flat porcelain or glass plates at a temperature of 100°. Remove the dry salt in flakes, and keep it in a stoppered bottle." Long-continued exposure to sunlight causes opacity in the scales, and renders them difficultly soluble (Wood).

*Ferri et Strychniæ Citras*, U. S. P., is prepared by mixing a solution containing five grains of strychnia and five grains of citric acid with a solution containing five hundred grains of citrate of iron and

ammonium, evaporating the mixture at a temperature not exceeding 140° to a syrupy consistence, and scaling in the usual way by spreading it upon plates of glass.

*Ferrum Tartaratum*, B. P.; *Ferri et Potassii Tartras*, U. S. P.—Ferric hydrate is dissolved in solution of acid tartrate of potassium, and the whole evaporated to dryness.

The ferric hydrate obtainable from five and a half (six is better) fluidounces of the official solution of ferric sulphate by the action of ammonia, in the manner detailed in the previous paragraphs, is mixed (in a mortar), while still moist but well drained, with two ounces of acid tartrate of potassium. The whole is set aside for about twenty-four hours, occasionally triturated to promote contact and reaction of the molecules (otherwise somewhat sluggish in attacking each other), and then heated in a dish over a water-bath to a temperature not exceeding 140° F.; a pint of distilled water is then added, and the mixture kept warm until nothing more will dissolve; filtered, evaporated at a temperature not exceeding 140° (greater heat causes decomposition), and, when the mixture has the consistence of syrup, spread on panes of glass and allowed to dry (in any warm and light place shown by a thermometer to be not hotter than 140°). The dry salt is then obtained in flakes. It should be kept in well-closed bottles.

*Ferri et Ammonii Tartras*, U. S. P., is made by saturating solution of acid tartrate of ammonium with ferric hydrate, evaporating, and scaling. The acid tartrate is prepared by exactly neutralizing half of any quantity of tartaric acid by carbonate of ammonium, and then adding the other half.

*The foregoing* are the only official scale preparations of iron. Many others of similar character might be formed. The *Citrate* dissolves slowly in cold but readily in warm water. None crystallize or give other indications of definite chemical composition Their properties are only constant so long as they are made with unvarying proportions of constituents. Want of chemical compactness, the loose state in which the iron is combined, precludes their recognition as well-defined chemical compounds, yet possibly enables them to be more readily assimilated as medicines than some of the more definite ferrous and ferric salts.

A definite *ferrous tartrate* ($FeC_4H_4O_6$) and *ferrous citrate* ($FeHC_6H_5O_7,H_2O$) have been obtained by reaction of iron and acid in hot water. They occur as white masses of microscopic crystals.

*Wine of Iron*, or "Steel" wine (*Vinum Ferri*, B. P.), made by digesting iron wire in sherry wine, probably contains tartrate of potassium and iron and other iron salts, formed by action of the metal on the acid tartrate of potassium and tartaric, citric, malic, and acetic acid present in the wine. *Vinum Ferri Citratis*, B. P., is a solution of ammonio-citrate of iron in orange wine.

## Black Hydrate of Iron. Ferroso-ferric Hydrate.

*Ferri Oxidum Magneticum*, B. P.

*Fifteenth Synthetical Reaction.*—To two-thirds of a small quantity of a solution of ferrous sulphate add a little sulphuric acid; warm, and gradually add nitric acid, as described in the tenth reaction, care being taken not to allow one drop more nitric acid than necessary to fall into the test-tube. Add the other third of ferrous sulphate, shake, and pour the liquid into excess of an alkali; black (at first brown) hydrate of iron, or ferroso-ferric hydrate ($Fe_3 8HO = Fe2HO, Fe_2 6HO$), is produced.

$$\underset{\text{Ferric sulphate.}}{Fe_2 3SO_4} + \underset{\text{Ferrous sulphate.}}{FeSO_4} + \underset{\text{Soda.}}{8NaHO} = \underset{\text{Blk. hydrate of iron.}}{Fe_3 8HO} + \underset{\text{Sulphate of sodium.}}{4Na_2SO_4}$$

It is so readily attracted by a magnet, even when moist, as to collect round the latter when immersed in the supernatant liquid. Hence the B. P. name, *Ferri Oxidum Magneticum.*

In this process the nitric acid oxidizes the hydrogen of the sulphuric acid, the sulphuric radical uniting with the ferrous sulphate, the iron of which is at the same time altered from the ferrous to the ferric condition, ferric sulphate being formed. If too much nitric acid be employed, the second portion of ferrous sulphate will also be converted into ferric salt, and the solution, on the addition of alkali, yield only red ferric hydrate. This result may be avoided by evaporating the solution of ferric sulphate nearly to dryness, thus boiling off excess of nitric acid, or by pouring first the ferric and then the ferrous liquid into the alkali and thoroughly stirring the mixture; any nitric acid is then neutralized and rendered incapable of oxidizing the ferrous sulphate subsequently added.

Black hydrate of iron is decomposed by heat, yielding, in a closed vessel, oxyhydrates, and, finally, black oxide of iron or ferroso-ferric oxide. Heated in the air it absorbs oxygen and gives ferric oxide. The black *forge-scales*, which collect near the blacksmith's anvil, have the composition of ferroso-ferric oxide: the black magma formed on exposing a mixture of iron and water to the air is ferroso-ferric hydrate; but these varieties are apt to contain particles of metal, and hence give hydrogen gas when dissolved in acids—a character which distinguishes them from the official preparation.

If a dried specimen of the black hydrate of iron be required, the mixture should be well boiled and then set aside for an hour or two to favor aggregation of the particles, the mixture filtered, and the precipitate washed until the washings contain no trace of sulphate (indicated by a white precipitate with chloride of barium). Black hydrate of iron absorbs oxygen even at the temperature of the water-bath; it should consequently be dried at 120°, a temperature at which only slight oxidation occurs.

## Pernitrate of Iron. Ferric Nitrate.

*Sixteenth Synthetical Reaction.*—Place a few iron tacks in dilute nitric acid and set aside; solution of ferric nitrate, or pernitrate of iron, is formed ($Fe_2 6NO_3$).

$$\underset{\text{Iron.}}{Fe_2} + \underset{\text{Nitric acid.}}{8HNO_3} = \underset{\text{Ferric nitrate.}}{Fe_2 6NO_3} + \underset{\text{Water.}}{4H_2O} + \underset{\text{Nitric oxide.}}{2NO}$$

This solution, made with care, and of a prescribed strength, forms the *Liquor Ferri Pernitratis*, B. P. (sp. gr. 1.107) and U. S. P. (sp. gr. 1.065). The process is as follows: four and a half fluidounces of nitric acid are diluted with sixteen ounces of distilled water, and one ounce of iron wire, free from rust, dissolved in the mixture, the latter being kept cool, to avoid violence of action. The liquid is finally filtered and diluted to thirty fluidounces.

Ferric nitrate and ferric acetate unite to form various aceto-nitrates, amongst which is one having the formula $Fe_2 4C_2H_3O_2NO_3HO,4H_2O$, crystallizing in hard, shining, brownish-red prisms.

## Reduced Iron.

*Seventeenth Synthetical Reaction.*—Pass hydrogen gas (dried by passing over pieces of chloride of calcium contained in a

Fig. 32.

Preparation of Reduced Iron.

tube, or through sulphuric acid in a wash bottle) into a small quantity of ferric oxyhydrate or oxide ("subcarbonate," U. S. P.) contained in a tube arranged horizontally (a test-tube, the bottom of which has been accidentally broken, answers very well), the oxide being kept hot by a gas-flame; oxygen is removed from the oxide by the hydrogen, steam escapes at the open end of the tube, and after a short time, when moisture

ceases to be evolved, metallic iron, in a minute state of division, remains. (See Fig. 32.)

$$\underset{\text{Ferric oxide.}}{Fe_2O_3} + \underset{\text{Hydrogen.}}{3H_2} = \underset{\text{Iron.}}{Fe_2} + \underset{\text{Water.}}{3H_2O}$$

While still hot throw the iron out into the air; it takes fire and falls to the ground as oxide.

If the ferric oxide is reduced in a gun-barrel heated by a strong furnace, the particles of iron aggregate to some extent, and, when cold, are only slowly oxidized in dry air. This latter form of reduced iron is *Fer réduit* or *Quevenne's Iron*, *the ferri pulvis*, or *Ferrum Redactum*, B. P. and U. S. P.—"a fine grayish-black powder, strongly attracted by the magnet, and exhibiting metallic streaks when rubbed with firm pressure in a mortar." It is often administered in the form of lozenges (*Trochisci Ferri Redacti*, B. P.), gum and sugar protecting the iron from oxidation as well as forming a vehicle for its administration.

*Note* I.—The spontaneous ignition of the iron in the above experiment is an illustration of the influence of minute division on chemical affinity. The action is the same as occurs whenever iron rusts, and the heat evolved and amount of oxide formed is not greater from a given quantity of iron; but the surface exposed to the action of the oxygen of the air is, in the case of this variety of reduced iron, so enormous compared with the weight of the iron, that heat cannot be conducted away sufficiently fast to prevent elevation of temperature to a point at which the whole becomes incandescent. In the slow rusting of iron, escape of heat occurs, but is not observed, because spread over a length of time; in the spontaneous ignition of reduced iron the whole is evolved at one moment. The mixture of lead and carbon (lead pyrophorus) resulting when tartrate of lead is heated in a test-tube until fumes cease to be evolved, spontaneously ignites when thrown into the air, and for the same reason. Many substances, solid and liquid, if sufficiently finely divided and liable to oxidation, and especially if exposed in a warm place, become hot and even occasionally spontaneously burst into flame. Oil on cotton-waste, powdered charcoal, coal, especially if pyritic or if very porous, or if powdered, resins in powder, and even flour, are familiar illustrations of materials liable to "heat" or even burn spontaneously.

*Note* II.—The student having time and opportunity for the experiment is advised to make this seventeenth reaction a roughly quantitative one, by way of realizing what has been stated (see, again, the General Principles of Chemical Philosophy, pp. 36–60) respecting the action of chemical force on definite weights only of matter. Three tubes, similar to the oxide-tube shown in the engraving, should be prepared, the second being connected to the first and the third to the second by India-rubber tubing in the usual manner. The first tube should contain pieces of chloride of calcium to absorb any traces of moisture not retained by the sulphuric acid. The

second tube (the ends of the small tube being temporarily closed by small corks) should be weighed in any ordinary scales which will turn with a quarter or half of a grain, and, the weight being noted, 160 grains of dry ferric oxide should be neatly placed in the middle of the tube. (The oxide must be previously gently heated in a small crucible over a lamp to remove all traces of moisture.) The third tube should contain pieces of chloride of calcium to absorb the water produced in the reaction, and just before being connected should be weighed. The operation is now carried out. At its close, and when the middle tube is cold, the latter tube and the third tube are again weighed. The oxide-tube should weigh 48 grains less than before, and the terminal tube 54 grains more than before.

$$\underbrace{Fe_2O_3}_{112+48=160} + \underbrace{3H_2}_{6} = \underbrace{Fe_2}_{112} + \underbrace{3H_2O}_{54}$$

The operation is more quickly and easily performed if one-half or one-quarter of the weight of oxide be taken; in that case one-half or one-quarter of the weight of iron and of water will be obtained. Indeed any weight of oxide may be employed; the amount of iron and water resulting will be *always exactly proportionate* to the weights just mentioned. Thus 16 parts of oxide yield 11.2 of iron and 5.4 of water. Iron, hydrogen, and oxygen always combine in proportions of 56, 1, and 16 respectively; hence our justification for agreeing that the symbol Fe shall stand for 56 parts by weight of iron, H for 1 part by weight of hydrogen, and O for 16 parts by weight of oxygen.

### Ferric Pyrophosphate.

*Eighteenth Synthetical Reaction.*—To solution of pyrophosphate of sodium add solution of persulphate of iron; a yellowish-white precipitate of ferric pyrophosphate ($Fe_4 3P_2O_7, 9H_2O$) separates. This precipitate, dissolved in solution of citrate of ammonium, and evaporated, yields apple-green scales (*Ferri Pyrophosphas*, U. S. P., containing forty-eight per cent. of anhydrous pyrophosphate).

### (*b*) *Reactions having Analytical Interest* (*Tests*).

(The iron occurring as a ferrous salt.)

*First Analytical Reaction.*—Pass sulphuretted hydrogen ($H_2S$) through a solution of a ferrous salt (*e. g.*, ferrous sulphate) slightly acidulated by hrodrochloric acid; no precipitate occurs.

This is a valuable negative fact, as will be evident presently.

*Second Analytical Reaction.*—Add sulphydrate of ammonium ($NH_4HS$) to solution of a ferrous salt; a black precipitate of ferrous sulphide (FeS) falls.

$$FeSO_4 + 2AmHS = FeS + Am_2SO_4 + H_2S.$$

*Third Analytical Reaction.*—Add solution of ferrocyanide of potassium (yellow prussiate of potash), $K_4Fe''Cy_6$, or $K_2Fcy''''$, to solution of a ferrous salt; a precipitate ($K_2Fe''Fe''Cy_6$, or $K_2Fe''Fcy$) falls, at first white, or bluish-gray, but rapidly becoming blue, owing to absorption of oxygen.

*Fourth Analytical Reaction.*—To solution of a ferrous salt add ferridcyanide of potassium (red prussiate of potash), $K_6Fe'''_2Cy_{12}$, or $K_6Fdcy$; a precipitate ($Fe''_3Fe'''_2Cy_{12}$, or $Fe''_3Fdcy$) resembling Prussian blue (Turnbull's blue) is thrown down.

*Other Analytical Reactions.*—The precipitates produced from ferrous solutions on the addition of alkaline carbonates, phosphates, and arseniates, as already described in the synthetical reactions of ferrous salts, are characteristic, and hence have a certain amount of analytical interest, but are inferior in this respect to the four reactions above mentioned.

*Note.*—Alkalies (potash, soda, or ammonia) are incomplete precipitants of ferrous salts, hence are almost useless as tests. To solution of a ferrous salt add ammonia ($NH_4HO$); on filtering off the whitish ferrous hydrate and testing the solution with sulphydrate of ammonium, iron will still be found. To another portion of the ferrous solution add a few drops of nitric acid and boil; this converts the ferrous into ferric salt, and now alkalies will wholly remove the iron, as already twice seen during the performance of the synthetical experiments.

In actual analysis, the separation of iron as ferric hydrate is an operation of frequent performance. This is always accomplished by the addition of alkali, and, if the iron occurs as a ferrous salt, by previous ebullition with a little nitric acid. Ferrocyanide and ferridcyanide of potassium are the reagents used in distinguishing ferrous from ferric salts.

(The iron occurring as a ferric salt.)

*Sixth Analytical Reaction.*—Through a ferric solution (ferric chloride *e. g.*) pass sulphuretted hydrogen; a white precipitate of the sulphur of the sulphuretted hydrogen falls, and the ferric is reduced to a ferrous salt, the latter remaining in solution. This reaction is of frequent occurrence in practical analysis.

$$2Fe_2Cl_6 + 2H_2S = 4FeCl_2 + 4HCl + S_2.$$

*Seventh Analytical Reaction.*—Add sulphydrate of ammonium to a ferric solution; the latter is reduced to the ferrous state, and black ferrous sulphide (FeS) is precipitated as in the second analytical reaction, sulphur being set free.

*Eighth Analytical Reaction.*—To a ferric solution add ferrocyanide of potassium ($K_4FeCy_6$, or $K_4Fcy''''$); a precipitate of Prussian blue, the common pigment, occurs ($Fe'''_4 3Fe''Cy_6$, or $Fe'''_4Fcy''''_3$). (*Ferri Ferrocyanidum*, U. S. P.)

*Ninth Analytical Reaction.*—To a ferric solution add solution of ferridcyanide of potassium; no precipitate occurs, but the liquid is darkened to a greenish or olive hue, according to the strength.

*Tenth Analytical Reaction.*—This is the production of a red precipitate of ferric hydrate, on the addition of alkalies to ferric salts, and is identical with the twelfth synthetical reaction.

*Note.*—This reaction illustrates the conventional character of the terms synthesis and analysis. It is of equal importance to the manufacturer and the analyst, and is synthetical or analytical according to the intention with which it is performed.

*Other ferric reactions* have occasional analytical interest. In neutral ferric solutions the tannic acid in aqueous infusion of galls occasions a bluish-black inky precipitate, the basis of most black writing inks.——(The *Mistura Ferri Aromatica* of the British Pharmacopœia, made by digesting metallic iron in an infusion of various vegetable substances, contains tannate, or rather tannates of iron: it is commonly known in Ireland by the name of Heberden's Ink, after the physician by whom it was first used. It contains about 1 grain of iron in 1 pint.) ——*Sulphocyanide of Potassium* (KCyS) causes the formation of ferric sulphocyanate, which is of deep blood-red color. ——There is no ferric carbonate; alkaline carbonates cause the precipitation of ferric hydrate, while carbonic acid gas escapes.

*Note.*—Cyanogen (CN, or Cy'), ferrocyanogen ($FeC_6N_6$, or $FeCy_6$, or simply $Fcy''''$), and ferridcyanogen ($Fe_2Cy_{12}$, or $Fdcy^{VI}$), are radicals which play the part of non-metallic elements, just as ammonium in its chemical relations resembles the metallic elements. They will be again referred to.

*Memorandum.*—The reader must on no account omit to write out equations or diagrams expressive of each of the reactions of iron, analytical as well as synthetical. It is presumed that this has already been done immediately after each reaction has been performed.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS CONTAINING **ONE** OF THE METALS, ZINC, ALUMINIUM, IRON.

Add solution of ammonia gradually:—

A dirty-green precipitate indicates iron in the state of a ferrous salt.

A red precipitate indicates iron in the state of a ferric salt.

A white precipitate, insoluble in excess, indicates the presence of an aluminium salt.

A white precipitate, soluble in excess, indicates zinc.

These results may be confirmed by the application of some of the other tests to fresh portions of the solution.

TABLE OF SHORT DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTION TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF **ONE, TWO, OR ALL THREE** OF THE METALS, ZINC, ALUMINIUM, IRON.

Boil about half a test-tubeful of the solution with a few drops of nitric acid. This insures the conversion of ferrous into ferric salts and enables the next reagent (ammonia) *completely* to precipitate the iron. Add excess of ammonia, and shake the mixture. Filter.

<table>
<tr><td colspan="2">Precipitate<br>Al Fe.*<br>Dissolve in HCl, add excess of KHO, stir, filter.</td><td>Filtrate<br>Zn.<br>Test by AmHS<br>(white ppt.).</td></tr>
<tr><td>Ppt.<br>Fe<br>(red ppt.).</td><td>Filtrate<br>Al.<br>Make slightly acid by HCl, and add excess of AmHO†<br>(white ppt.).</td><td></td></tr>
</table>

* The aluminium precipitate ($Al_2 6HO$) is white, the iron ($Fe_2 6HO$) red. If the precipitate is red, iron must be and aluminium may be present; if white, iron is absent, and further operation on the precipitate unnecessary. This precipitate ($Al_2 6HO$ and $Fe_2 6HO$) may also, if sufficient is at disposal, be analyzed by simply well shaking a washed portion in a tube with solution of potash or soda; the hydrate of iron is not thereby affected, while the hydrate of aluminium is dissolved, and may be detected in the clear decanted fluid by neutralizing all alkali by a little excess of acid, and then adding excess of ammonia.

† Alumina, when in small quantity, is sometimes prevented from being precipitated by ammonia through the presence of organic mat-

*Note* I.—If iron is present, portions of the original solution must be tested by ferridcyanide of potassium for ferrous, and by ferrocyanide for ferric salts; dark-blue precipitates with both indicate both salts.

*Note* II.—If no ferrous salt is present, ebullition with nitric acid is unnecessary. It is, perhaps, therefore advisable always to determine this point by previously testing a little of the original solution with ferridcyanide; if no blue precipitate occurs, the nitric acid treatment may be omitted.

### Chart for all Metals hitherto considered.

The following Table (*vide* p. 164) is perhaps the best, but not the only adaptation of the ordinary reactions to systematic analysis. It is little else than the addition of the analytical scheme for the third group to that of the first two groups. As before, analysis is commenced by the addition of chloride of ammonium ($NH_4Cl$) to prevent partial precipitation of magnesium, and ammonia ($NH_4HO$) to neutralize any acid. The latter would attack the chief group-precipitant, sulphydrate of ammonium ($NH_4HS$), preventing its useful action, and causing a precipitation of the sulphur it commonly contains.

*Note.*—When a test gives no reaction, absence of the body sought for may be fairly inferred. If a group-test (that is, a test which precipitates a group of substances) gives no reaction, the analyst is saved the trouble of looking for any of the members of that group.

---

ter derived from the filter paper by action of the potash. In cases of doubt, therefore, before adding ammonia boil the liquid with a little nitric acid, which destroys any organic matter.

TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF **ANY OR ALL** OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.

Add AmCl; AmHO; AmHS; stir, filter.

<table>
<tr><td colspan="3">Precipitate<br>Fe Al Zn.<br>Wash, dissolve in HCl,* boil (to remove $H_2S$), filter (to remove S), add excess of KHO,† stir, filter.</td><td colspan="4">Filtrate<br>Ba Ca Mg Am Na K.<br>Add $Am_2CO_3$, boil, filter.</td></tr>
<tr><td>Ppt.<br>Fe<br>(test orig. sol. by $K_4Fcy$ and $K_6Fdcy$).</td><td colspan="2">Filtrate<br>Al Zn.<br>Neutralize with HCl.<br>Add AmHO, stir, filter.</td><td colspan="2">Ppt.<br>Ba Ca.<br>Dissolve in $HC_2H_3O_2$,<br>Add $K_2CrO_4$, filter.</td><td colspan="2">Filtrate<br>Mg Am Na K.<br>Add $Am_2HPO_4$, stir, filter.</td></tr>
<tr><td></td><td>Ppt.<br>Al<br>(white).</td><td>Filtrate<br>Zn.<br>Add AmHS<br>(white ppt.).</td><td>Ppt.<br>Ba<br>(yellow).</td><td>Filtrate<br>Ca.<br>Add $Am_2C_2O_4$<br>(white ppt.).</td><td>Ppt.<br>Mg<br>(white).</td><td>Filtrate<br>Am Na K.<br>Evap., ignite, dissolve.<br>Na by flame; K by $PtCl_4$;<br>orig. sol. for Am.</td></tr>
</table>

* Add, also, a few drops of $HNO_3$ if Fe be present—*i. e.*, if the ppt. be black. (*Vide* notes on pp. 160 and 163.)

† Or add excess of ammonia, filter, and test filtrate for zinc. The hydrates of iron and aluminium may then be separated by an alkali, as described in the foot-note to the previous table, p. 162.

## QUESTIONS AND EXERCISES.

207. Name the chief ores of iron.

208. How is the metal obtained from the ores?

209. What is the chemical difference between cast iron, wrought iron, and steel?

210. Explain the process of welding.

211. What is the nature of chalybeate waters?

212. Illustrate by formula the difference between ferrous and ferric salts.

213. Under what different circumstances may the atom of iron be considered to exert bivalent, trivalent, and sexivalent activity?

214. Write a paragraph on the nomenclature of iron salts.

215. Give a diagram of the official process for the preparation of ferrous sulphate.

216. In what respects do Sulphate of Iron, Granulated Sulphate of Iron, and dried Sulphate of Iron differ?

217. How is ferrous sulphate obtained on the large scale?

218. Mention the chemical names of white, green, and blue vitriol.

219. Why does ferrous sulphate become brown by prolonged exposure to air?

220. Give a diagram showing the formation of Ferrous Carbonate.

221. Describe the action of atmospheric oxygen on ferrous carbonate. Can the effect be prevented?

222. In what order would you mix the ingredients of *Mistura Ferri Composita*, and why?

223. Write out an equation illustrative of the formation of the Phosphate of Iron.

224. Why is acetate of sodium used in the preparation of ferrous phosphate?

225. Which four compounds of iron may be formed by the direct union of their elements?

226. Give the official method for the preparation of Solution of Ferric Chloride.

227. Of what use is the spirit in Tincture of Perchloride of Iron?

228. How may Ferrous be converted into Ferric Sulphate?

229. What is the formula of Ferric Acetate? And how is it prepared for use in pharmacy?

230. Express by formulæ the difference between *Ferri Peroxidum Humidum*, B. P., and *Ferri Peroxidum Hydratum*, B. P.

231. How does Ferric Hydrate act as an antidote to arsenic?

232. What are the properties of anhydrous ferric oxide?

233. What are the general characters and mode of production of the medicinal scale preparations of iron?

234. In what state is the iron in *Vinum Ferri*, B. P.?

235. What other form of Wine of Iron is official in Great Britain?

236. Give equations illustrating the chief steps in the artificial production of the so-called Magnetic Oxide of Iron.

237. How is precipitated magnetic oxide of iron distinguished from the varieties made directly from the metal?

238. Why is magnetic oxide of iron officially directed to be dried at a temperature not exceeding 120° Fahr.?

239. Give a diagram showing the formation of Ferric Nitrate.

240. Work out a sum showing how much anhydrous ferric oxide will yield, theoretically, one hundred-weight of iron. Ans. 160 lbs.

241. What are the properties of anhydrous ferric oxide?

242. Give the characteristic tests for iron, distinguishing between ferrous and ferric reactions, and illustrating each by an equation or a diagram:—

*a*. Sulphydrate of ammonium.
*b*. Ferrocyanide of potassium.
*c*. Ferridcyanide of potassium.
*d*. Caustic alkalies.
*e*. Sulphocyanate of potassium.

243. Describe the action of ammonia on salts of iron, aluminium, and zinc respectively.

244. What precautions must be used in testing for calcium in the presence of iron?

245. How is magnesium detected in the presence of zinc?

246. How is aluminium detected in the presence of magnesium?

247. Draw up a scheme for the analysis of an aqueous liquid containing salts of iron, barium, and potassium.

248. How may zinc, magnesium, and ammonium be consecutively removed from aqueous solution?

Arsenicum. Sb.

## ARSENICUM, ANTIMONY.

These two elements resemble metals in appearance and in the character of some of their compounds; but they are still more closely allied to the non-metals, especially to phosphorus and nitrogen. They are quinquivalent ($As^{v}$, $Sb^{v}$), as seen in arsenic anhydride ($As_2O_5$) and pentachloride of antimony ($SbCl_5$), but usually exert trivalent activity only ($As^{III}$, $Sb^{III}$), as seen in the hydrogen and other compounds ($AsH_3$, $AsCl_3$, $AsBr_3$, $AsI_3$). A few preparations of these elements are used in medicine; but all are more or less powerful poisons, and hence have considerable toxicological interest. The iodide (*Arsenici Iodidum*, U. S. P.) is made by cautiously fusing together atomic proportions of arsenicum and iodine. It is an orange-red crystalline solid soluble in water. The *Liquor Arsenici et Hydrargyri Iodidi*, U. S. P., is made by dissolving iodide of arsenicum and red iodide of mercury in water, in the proportion of 1 grain of each of the solids to 204 grains of water.

Arsenicum is an exception to the rule that the atomic weights (taken in grains, grammes, or other weight) of elements, under similar circumstances of temperature and pressure, give equal volumes of vapor, the equivalent weight (75) of arsenicum only occupying half such a volume. Hence, while the molecular weights (that is, double the atomic weights of oxygen ($O_2 = 32$), hydrogen ($H_2 = 2$), nitrogen ($N_2 = 28$), etc., give a similar bulk of vapor at any given temperature and pressure, the double atomic weight of arsenicum

($As_2$=150), at the same temperature and pressure, only affords half this bulk. It would appear, therefore, that the molecule of arsenicum contains four atoms, and that its formula is $As_4$. As in the case of sulphur, however, arsenicum, in the state ordinarily known to us, may be abnormal, and a variety yet be found in which the molecular weight is double (instead of quadruple) the atomic weight.

From observed analogy between the two metals, the molecular constitution of antimony is probably similar to that of arsenicum.

---

## ARSENICUM.

Symbol As. Atomic weight 75.

*Sources.*—Arsenical ores are frequently met with in nature, the commonest being the arsenio-sulphide of iron (FeSAs). This mineral is roasted in a current of air, the oxygen of which, combining with the arsenicum, forms common white arsenic ($As_2O_3$) (*Acidum Arsenicosum*, B. P. and U. S. P.), which is condensed in chambers or long flues. It commonly "occurs as a heavy white powder, or in sublimed masses, which usually present a stratified appearance, caused by the existence of separate layers, differing from each other in degrees of opacity." The vitreous or amorphous arsenic is far more soluble than the crystalline variety, and in other respects they differ in properties. Such differences between the crystalline and amorphous varieties of an element or compound are not unfrequent: they have not yet been satisfactorily explained. *Realgar* (red algar) is the red native sulphide ($As_2S_2$), and orpiment (*auripigmentum*, the golden pigment) the yellow native sulphide ($As_2S_3$) of arsenicum. The iodide of arsenicum ($AsI_3$) may be made from its elements or (Babcock) by dissolving white arsenic in aqueous hydriodic acid and evaporating.

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Reactions having Synthetical Interest.*

#### Alkaline Solution of Arsenic.

*First Synthetical Reaction.*—Boil a grain or two of powdered arsenic ($As_2O_3$) in water containing an equal weight of carbonate of potassium, and, if necessary, filter. The solution, colored with compound tincture of lavender, and containing 4 grains of arsenic per ounce, forms the *Liquor Arsenicalis*, B. P., or *Liquor Potassii Arsenitis*, U. S. P. (*Fowler's Solution*).

*Note.*—This official solution does not generally contain arsenite of potassium; for the arsenic does not decompose the carbonate of potassium, or only after long boiling. From concentrated solutions carbonic acid gas is more quickly eliminated.

## Arsenious Acids and other Arsenites.

Arsenic or *arsenious anhydride* (the so-called arsenious acid), when dissolved in water, is said to yield true arsenious acid ($H_3AsO_3$) —the arsenite of hydrogen.

$$\underset{\text{Arsenious anhydride.}}{As_2O_3} + \underset{\text{Water.}}{3H_2O} = \underset{\text{Arsenious acid.}}{2H_3AsO_3}$$

When arsenic ($As_2O_3$) is dissolved in excess of solutions of potash or soda, arsenites are formed having the formulæ $KH_2AsO_3$ and $NaH_2AsO_3$. Boiled with excess of arsenic, one molecule of these salts combines with one of arsenic. The usual character of such compounds is that of oily alkaline liquids. Arsenic fused with alkaline carbonates yields pyroarseniates ($Na_4As_2O_7$ or $H_4As_2O_7$) and metallic arsenicum. Arsenites have the general formula $R'_3AsO_3$.

## Acid Solution of Arsenic.

*Second Synthetical Reaction.*—Boil arsenic with dilute hydrochloric acid. Such a solution made with prescribed proportions of acid and water, and containing 4 grains of arsenic ($As_2O_3$) per ounce, forms the *Liquor Arsenici Hydrochloricus*, B. P. (*Liquor Arsenici Chloridi*, U .S. P.). (*De Valangin's Solution* contained a grain and a half per ounce.)

*Note.*—No decomposition occurs in this experiment. The liquid is simply a solution of arsenic in dilute hydrochloric acid. These two solutions may be preserved for analytical operations.

*Mem.*—The practical student should boil arsenic in water also, and thus have an acid, alkaline, and aqueous solution for analytical comparison.

## Arsenicum.

*Third Synthetical Reaction.*—Place a grain or less of arsenic at the bottom of a narrow test-tube, cover it with about half an inch or an inch of small fragments of dry charcoal, and hold the tube, nearly horizontally, in a flame, the mouth being loosely covered by the thumb. At first let the bottom of the tube project slightly beyond the flame, so that the charcoal may become nearly red hot; then heat the bottom of the tube. The arsenic will sublime, become deoxidized by the charcoal, carbonic oxide being formed, and arsenicum deposited in the cool part of the tube as a dark mirror-like incrustation.

There is a characteristic odor, resembling garlic, emitted during this operation, probably due to a partially oxidized trace of arsenicum, which escapes from the tube; for arsenic does not give this odor; moreover, arsenicum being a freely oxidizable element, its

vaporous particles could scarcely exist in the air in an entirely unoxidized state.

*Metallic arsenicum* (*Arsenicum*, U. S. P.) may be obtained in large quantities by the above process if the operation be conducted in vessels of commensurate size. But performed with great care, in narrow tubes, using not charcoal alone, but *black flux* (a mixture of charcoal and carbonate of potassium obtained by heating acid tartrate of potassium in a test-tube or other closed vessel till no more fumes are evolved), the reaction has considerable analytical interest, the garlic odor and the formation of the mirror-like ring being highly characteristic of arsenicum. Compounds of mercury and antimony, however, give sublimates which may be mistaken for arsenicum.

## Arsenic Acid and other Arseniates.

*Fourth Synthetical Reaction.*—Boil a grain or two of arsenic with a few drops of nitric acid until red fumes cease to be evolved; evaporate the solution in a small dish to dryness, to remove excess of nitric acid; dissolve the residue in water; the product is Arsen'ic acid ($H_3AsO_4$).

Arsenic acid, when strongly heated, loses the elements of water, and arsenic anhydride remains ($As_2O_5$).

Arsenic anhydride readily absorbs water and becomes arsenic acid ($H_3AsO_4$). Arsenic acid is readily reduced to arsenious by the action of reducing agents such as sulphurous acid, $H_3AsO_4 + H_2SO_3 = H_3AsO_3 + H_2SO_4$, or oxalic acid ($H_3AsO_4 + H_2C_2O_4 = H_3AsO_3 + H_2O + 2CO_2$).

Salts analogous to arsenic acid, the arseniate of hydrogen, are termed *arseniates*, and have the general formula $R'_3AsO_4$. The ammonium arseniate ($Am_2HAsO_4$) may be made by neutralizing arsenic acid with ammonia. Its solution in water forms a useful reagent. ——Arsenic acid is used as an oxidizing agent in the manufacture of the well-known dye, magenta.

Arsenite and arseniate of sodium are used in the cleansing operations of the calico-printer.

## Pyroarseniate and Arseniate of Sodium.

*Fifth Synthetical Reaction.*—Fuse two or three grains of common white arsenic ($As_2O_3$) with nitrate of sodium ($NaNO_3$) and dried carbonate of sodium ($Na_2CO_3$) in a porcelain crucible, and dissolve the mass in water; solution of arseniate of sodium ($Na_2HAsO_4$) results.

$$\underset{\text{Arsenic.}}{As_2O_3} + \underset{\substack{\text{Nitrate of}\\\text{sodium.}}}{2NaNO_3} + \underset{\substack{\text{Carbonate}\\\text{of sodium.}}}{Na_2CO_3} = \underset{\substack{\text{Pyroarseniate}\\\text{of sodium.}}}{Na_4As_2O_7} + \underset{\substack{\text{Nitrous}\\\text{anhydride.}}}{N_2O_3} + \underset{\substack{\text{Carbonic}\\\text{acid gas.}}}{CO_2}$$

The official proportions (B. P.) are 10 of arsenic to 8½ of nitrate of sodium and 5½ of dried carbonate, each powdered, the whole well mixed, fused in a crucible at a red heat till effervescence ceases, and

the liquid poured out on a slab. The product is pyroarseniate of sodium ($Na_4As_2O_7$). Dissolved in water, crystallized and dried, the salt has the formula $Na_2HAsO_4,7H_2O$ (*Sodæ Arsenias*, B. P.; *Sodii Arsenias*, U. S. P.).

$$Na_4As_2O_7 + 15H_2O = 2(Na_2HAsO_4,7H_2O).$$

Heated to 300° F. the crystals lose all water. A solution of 4 grains of the anhydrous salt ($Na_2HAsO_4$) in 1 ounce of water forms the *Liquor Sodii Arseniatis*, U. S. P. The anhydrous salt is used in this preparation because the crystallized is of somewhat uncertain composition. The fresh crystals are represented by the formula $Na_2HAsO_4,12H_2O$ (=53.7 per cent. of water); these soon effloresce and yield a stable salt having the formula $Na_2HAsO_4,7H_2O$ (= 40.4 per cent. of water). To avoid the possible employment of a mixture of these bodies, the invariable anhydrous salt is officially used, constancy in the strength of a powerful preparation being thereby secured.

The student will find useful practice in verifying the above numbers representing the centesimal proportion of water in the two arseniates of sodium. This will readily be accomplished if what has already been stated respecting a symbol representing a number as well as a name, and the remarks concerning a molecular weight, be remembered.

The shape of each of the two varieties of arseniate of sodium ($Na_2HAsO_4,12H_2O$, and $Na_2HAsO_4,7H_2O$) is identical with that of the corresponding phosphate of sodium ($Na_2HPO_4,12H_2O$, and $Na_2HPO_4,7H_2O$); the structure of the molecule of the 12-arseniate is the same as that of the 12-phosphate, and the 7-arseniate as that of the 7-phosphate; the two former are isomorphous, the two latter are isomorphous. This is only one instance of the strong analogy of arsenicum and its compounds with phosphorus and its corresponding compounds. The preparation and characters of the next substance, arseniate of iron, will remind the learner of phosphate of iron.

## Arseniate of Iron. Ferrous Arseniate.

*Sixth Synthetical Reaction.* — To solution of arseniate of sodium add a little acetate of sodium and then solution of ferrous sulphate, a precipitate of ferrous arseniate occurs ($Fe_2AsO_4$) (*Ferri Arsenias*, B. P.). On the large scale 4 parts of dried arseniate and 3 of acetate dissolved in 40 of water, mixed with 9 of sulphate in 60 of water, may be employed. The precipitate should be collected on a calico filter, washed, squeezed, and dried at a low temperature (100° F.) over a water-bath to avoid excessive oxidation.

$$\underset{\text{Arseniate of sodium.}}{2Na_2HAsO_4} + \underset{\text{Acetate of sodium.}}{2NaC_2H_3O_2} + \underset{\text{Ferrous sulphate.}}{3FeSO_4} = \underset{\text{Ferrous arseniate.}}{Fe_2AsO_4} + \underset{\text{Sulphate of sodium.}}{3Na_2SO_4} + \underset{\text{Acetic acid.}}{2HC_2H_3O_2}$$

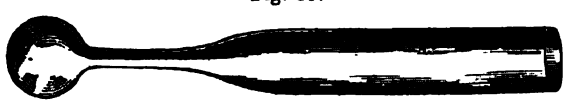
*The use of the acetate of sodium* is to insure the absence of free sulphuric acid in solution; sulphate of sodium being formed together with acetic acid. Sulphuric acid is a solvent of ferrous arseniate; acetic acid is not. It is impossible to prevent the separation of sulphuric acid, if only ferrous sulphate and arseniate of sodium be employed. At the instant of precipitation ferrous arseniate is white, but rapidly becomes of a green or greenish-blue color, owing to absorption of oxygen and formation of a ferroso-ferric arseniate. It is a tasteless amorphous powder, soluble in acids.

The *Hydride* and *Sulphides* of *Arsenicum*, and the *Arsenites* and *Arseniates of Copper* and of *Silver* are mentioned in the following analytical paragraphs:—

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—Repeat the third synthetical reaction, operating on not more arsenic than has about the bulk of a small pin's head, and using not charcoal alone, but *black flux* (a mixture of charcoal and carbonate of potassium obtained by heating acid tartrate of potassium in a test-tube or other nearly closed vessel till fumes cease to be evolved). The tube employed should be a narrow test-tube, or, better, a tube (easily made from glass tubing) having the following (Berzelius's) form:—

Fig. 33.



The arsenic and black flux are placed in the bulb of the tube, which is then heated in a flame; the arsenicum condenses on the constricted portion of the tube. If now the bulb be carefully fused off in a flame, the arsenicum may be chased up and down the narrower part of the tube until the air in the tube has reoxodized it to arsenious anhydride.

If the operation has been performed in a less delicate manner in an ordinary test-tube, cut or break off portions of the tube containing the sublimate of arsenicum, put them into a test-tube and heat the bottom of the latter, holding it nearly horizontally, and partially covering the mouth with the finger or thumb; the arsenicum ($As_4$) will absorb oxygen from the air in the tube, and the resulting arsenious anhydride ($As_2O_3$) be deposited on the cool part of the tube in brilliant transparent, generally imperfect, octahedral crystals.

*Microscopic Test.*—Prove that the crystals are identical in form with those of common white arsenic, by heating a grain or less of the latter in another test-tube, examining the two sublimates by a good lens or compound microscopic.

Fig. 34.

A Sublimate of White Arsenic. (Magnified.)

Fig. 34*a*.

A perfect Octahedron.

The appearance of a sublimate of arsenic is peculiar and quite characteristic. The primary form of each crystal is an octahedron (ὀκτώ, *okto*, eight; ἵδρα, *hedra*, side) (fig. 34*a*), or, rarely, a tetrahedron, and in a sublimate a few perfect octahedra are generally present. Usually, however, the crystals are modifications of octahedra such as are shown in fig. 34—which is drawn from actual sublimates.

*Second Analytical Reaction.*—Place a thin piece of copper, about a quarter inch wide and half inch long, in a solution of arsenic, acidified by hydrochloric acid, and boil (nitric acid must not be present, or the copper itself will be dissolved); arsenicum is deposited on the plate in a metallic condition, an equivalent portion of copper going into solution. Pour off the supernatant liquid from the copper, wash the latter once or twice with water, dry the piece of metal by holding in the fingers and passing through a flame, and finally place it at the bottom of a clean dry narrow test-tube or a Berzelius tube; sublime as described in the last reaction, again noticing the form of the resulting crystals.

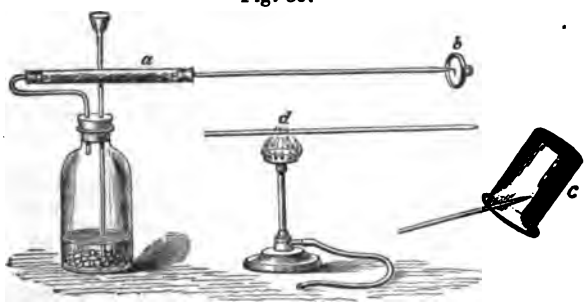
This is commonly known as Reinsch's test of arsenicum. The tube may be reserved for subsequent comparison with an antimonial sublimate (p. 185).

*Note.*—Copper itself frequently contains arsenicum, a fact that may not, perhaps, much trouble an operator so long as he is performing experiments in practical chemistry merely for educational purposes; but when he engages in the analysis of bodies of unknown

composition, he must assure himself that neither his apparatus nor materials already contain the element of which he is in search.

*The detection of arsenicum in metallic copper* is best accomplished by distilling a mixture of a few grains of the sample with five or six times its weight of ferric hydrate or chloride (free from arsenicum) and excess of hydrochloric acid. The arsenicum is thus volatilized in the form of chloride of arsenicum, and may be condensed in water and detected by sulphuretted hydrogen (5th Analytical Reaction) or Reinsch's test. The ferric chloride solution is, if necessary, freed from any trace of arsenicum by evaporating once or twice to dryness with excess of hydrochloric acid (Odling).

Fig. 35.




The Hydrogen Test for Arsenicum.

*Third Analytical Reaction.* — The hydrogen test, or "*Marsh's*" *test.*—Generate hydrogen in the usual way from water by zinc and sulphuric acid, a bottle of about four or six ounces capacity being used, and a funnel-tube and short delivery-tube passing through the cork in the usual manner (see above figure). Dry the escaping hydrogen (except in rough experiments, when it is unnecessary) by adapting to the delivery-tube, by a pierced cork, a short piece of wider tubing filled with fragments of chloride of calcium *a*. To the opposite end of the drying tube fit a piece of narrow tubing ten or twelve inches long, made of hard German glass, and having its aperture narrowed by drawing out in the flame of the blow-pipe. When the hydrogen has been escaping at such a rate and for a sufficient number of minutes as to warrant the operator in concluding that all the air originally existing in the bottle has been expelled, set light to the jet, and then pour eight or ten drops of the aqueous solution of arsenic; or three or four drops of the acid or alkaline solution of arsenic, previously prepared, into the funnel-tube, washing the liquid into the generating-bottle with a little water. The arsenic is at

once reduced to the state of arsenicum, and the latter combines with some of the hydrogen to form hydride of arsenicum or arseniuretted hydrogen gas ($AsH_3$). Immediately hold a piece of earthenware or porcelain (the lid of a porcelain crucible, *b*, if at hand) in the hydrogen jet at the extremity of the delivery-tube; a brown spot of arsenicum is deposited on the porcelain. Collect several of these spots, and retain them for future comparison with antimonial spots (p. 185.)

The separation of arsenicum in the flame is due to the decomposition of the arseniuretted hydrogen by the heat of combustion. The cool porcelain at once condenses the arsenicum, and thus prevents its oxidation to white arsenic, which would otherwise take place at the outer edge of the flame.

Hold a small beaker, *c*, or wide test-tube over the flame for a few minutes; a white film of arsenic ($As_2O_3$) will be slowly deposited, and may be further examined in contrast with a similar antimonial film (p. 185).

During these experiments the effect produced by the arsenical vapors on the color of the hydrogen-flame will have been noticed; they give it a dull livid bluish tint. This is characteristic.

Apply the flame of a gas-lamp to the middle of the hard delivery-tube, *d;* the arseniuretted hydrogen, as before, is decomposed by the heat, but the liberated arsenicum ($As_2$) immediately condenses in the cool part of the tube beyond the flame, forming a dark metallic mirror. The tube may be removed and kept for comparison with an antimonial deposit.

*Note* I.—Zinc, like copper, frequently itself contains arsenicum. When a specimen free from arsenicum is met with, it should be reserved for analytical experiments, or a quantity of guaranteed purity should be purchased of the chemical-apparatus maker. Sulphuric acid is more easily obtained free from arsenic.

*Note* II.—In delicate and important applications of Marsh's test, magnesium may be substituted for zinc with safety, as arsenicum has not yet been, and is not likely to be, found in magnesium. Magnesium in rods is convenient for this purpose, and may be obtained from most dealers in chemicals.

*Note* III.—Sulphuric acid, which is often used for drying gases, decomposes arseniuretted hydrogen. Chloride of calcium is therefore the appropriate desiccating agent for this gas.

*Fourth Analytical Reaction.—Fleitmann's test.*—Generate hydrogen by heating to near the boiling-point a strong solution of caustic soda or potash and some pieces of zinc.

($Zn + 2NaHO = H_2 + Na_2ZnO_2$—zincate of sodium.)

Drop into the test-tube a little arsenical solution, and spread over the mouth of the tube a cap of filter-paper moistened with one drop of solution of nitrate of silver. Again heat the tube, taking care that the liquid itself shall not spirt up on to the cap; the arsenic is reduced to arsenicum, the latter uniting with the hydrogen as in Marsh's test; and the arseniuretted hydrogen passing up through the cap reacts on the nitrate of silver, causing the production of a purplish-black spot.

$$AsH_3 + 3H_2O + 6AgNO_3 = H_3AsO_3 + 6HNO_3 + 3Ag_2.$$

*Note.*—This reaction is particularly valuable, enabling the analyst to quickly distinguish arsenicum in the presence of its sister element antimony, which, although it combines with the hydrogen evolved from dilute acid and zinc, does not combine with the hydrogen evolved from solution of alkali and zinc, and therefore does not give the effect just described.

*Fifth Analytical Reaction.*—To a solution of chloride of tin in strong hydrochloric acid add a very small quantity of any arsenical solution. Arsenicum then separates, especially on the application of heat, giving the mixture a yellowish and then brownish hue or grayish-brown turbidity, or even a sediment of gray-brown flocks, according to the amount present. Much water prevents the reaction; its presence, therefore, must be avoided as much as possible; indeed a liquid saturated by hydrochloric acid gas gives best results. Arsenic in sulphuric or hydrochloric acid or in tartar emetic, etc., may be detected by this method. Nitrates, such as subnitrate of bismuth, must first be heated with sulphuric acid to remove the nitric radical before applying this reduction test for arsenicum. The stannous is converted to stannic salt during the reaction.

---

*Distinction between Arsenious and Arsenic combinations.*—The above tests are those of arsenicum, whether existing in the arsenious or arsenic condition, though from the latter the element is not generally eliminated so quickly as from the former. Of the following reactions, that with nitrate of silver at once distinguishes arsenious acid and other arsenites from arsenic acid and other arseniates.

*Mem.*—The exact nature of all these analytical reactions will be more fully evident if traced out by diagrams or equations.

---

*Sixth Analytical Reaction.*—Through an acidified solution of arsenic pass sulphuretted hydrogen; a yellow precipitate of sulphide of arsenicum or arsenious sulphide ($As_2S_3$) quickly

falls. Add an alkaline hydrate or sulphydrate to a portion of the precipitate; it readily dissolves. The precipitate consequently would not be obtained on passing sulphuretted hydrogen through an alkaline solution of arsenic. To another portion of the precipitate, well drained, add *strong* hydrochloric acid; it is insoluble, unlike sulphide of antimony. Neither sulphide is soluble in the weak acid.

*Note* I.—Cadmium also affords a yellow sulphide in an acid solution by action of sulphuretted hydrogen, but this sulphide is insoluble in alkaline liquids. Under certain circumstances, tin, too, yields a yellow sulphide; but tin is otherwise easily distinguished (*vide* Tin).

*Note* II.—A trace of sulphide of arsenicum is sometimes met with in sulphur (distilled from arsenical pyrites). It may be detected by digesting the sulphur in solution of ammonia, filtering, and evaporating to dryness; a yellow residue of sulphide of arsenicum is obtained if that substance be present.

*Seventh Analytical Reaction.*—Through an acidified solution of arsenic acid, or any other arseniate, pass sulphuretted hydrogen; a yellow precipitate of arsenious sulphide and sulphur ($As_2S_3 + S_2$) slowly falls Both arsenious sulphide ($As_2S_3$) and arsenic sulphide ($As_2S_5$) are soluble in alkaline hydrates and sulphydrates.

*Chemical Analogy of Sulphur and Oxygen.*—The solubility of arsenious and arsenic sulphide in alkaline solutions is good evidence of the close chemical analogy between them and the corresponding oxygen compounds of arsenicum. The potassium arsenite and sulph-arsenite, arseniate and sulph-arseniate, have the composition represented by the following formulæ:—

$K_3AsO_3$ $\qquad$ $K_3AsO_4$
$K_3AsS_3$ $\qquad$ $K_3AsS_4$;

and the corresponding ammonium and sodium salts have a similar composition:—

$$6AmHS + As_2S_3 = 2Am_3AsS_3 + 3H_2S$$
$$6AmHS + As_2S_5 = 2Am_3AsS_4 + 3H_2S.$$

*Eighth Analytical Reaction.*—To an aqueous solution of arsenic add two or three drops of solution of sulphate of copper, and then cautiously add diluted solution of ammonia, drop by drop, until a green precipitate is obtained. The production of this precipitate is characteristic of arsenicum. To a portion of the mixture add an acid; the precipitate dissolves. To another portion add alkali; the precipitate dissolves. These two experiments show the advantage of testing a suspected arsenical solution by litmus-paper before applying this reaction; if acid, cautiously adding alkali, if alkaline, adding acid, till neutrality

is obtained. (Or a special copper reagent may be used; see a note to the Eleventh Analytical Reaction, below.)

The precipitate is arsenite of copper ($Cu''HAsO_3$) or *Scheele's Green*. More or less pure, or mixed with acetate or, occasionally, carbonate of copper, it is very largely used as a pigment under many names, such as Brunswick Green and Schweinfurth Green, by painters, paper-stainers, and others.

*Ninth Analytical Reaction.*—Apply the test just described to a solution of arsenic acid or other arseniate; a somewhat similar precipitate of arseniate of copper is obtained.

*Tenth Analytical Reaction.*—Repeat the eighth reaction, substituting nitrate of silver for sulphate of copper; in this case *yellow* arsenite of silver ($Ag_3AsO_3$) falls, also soluble in acids and alkalies.

*Eleventh Analytical Reaction.*—Apply the test to a solution of arsenic acid or other arseniate; a *chocolate*-colored precipitate of arseniate of silver ($Ag_3AsO_4$) falls.

*Copper and Silver Reagents for Arsenicum.*—The last four reactions may be performed with increased delicacy and certainty of result, if the copper and silver reagents be previously prepared in the following manner: To solution of pure sulphate of copper (about 1 part in 20 of water) add ammonia until the blue precipitate at first formed is nearly but not quite, redissolved; filter and preserve the liquid as an arsenicum reagent, labelling it *solution of ammonio-sulphate of copper* (B. P.). Treat solution of nitrate of silver (about 1 part in 40) in the same way, and label it *solution of ammonio-nitrate of silver* (B. P.). The composition of these two salts will be referred to subsequently.

*Arsenious and Arsenic Compounds.*—While many reagents may be used for the detection of arsenicum, only nitrate of silver, as already stated, will readily indicate in which state of oxidation the arsenicum exists; for the two sulphides and the two copper precipitates, though differing in composition, resemble each other in appearance, whereas the two silver precipitates differ in color as well as in composition.

Soluble arseniates also give insoluble arseniates with solutions of salts of barium, calcium, zinc, and some other metals.

In group-testing, arsenicum, if existing as arsenic acid or other arseniate, is not readily affected by such tests as sulphuretted hydrogen or even hydrogen itself. Hence, if its presence in that state is suspected, the liquid under analysis should be warmed with a little sulphurous acid or oxalic acid, and then tested with sulphuretted hydrogen.

*Antidote.*—In cases of poisoning by arsenic or arsenical preparations, the most effective antidote is recently precipitated moist ferric hydrate (*Ferri Peroxidum Humidum*, B. P.; *Ferri Oxidum Hydratum*, U. S. P.). It is perhaps best

administered in the form of a mixture or solution of perchloride of iron (*Liquor* or *Tinctura*) with carbonate of sodium—two or three ounces of the former to about one ounce of the crystals of the latter. Instead of the carbonate of sodium, about a quarter of an ounce of calcined magnesia may be used. These quantities will render at least 10 grains of arsenic insoluble. Emetics should also be given, and the stomach pump applied as quickly as possible.

The above statements regarding the antidote for arsenic may be verified by mixing the various substances together, filtering, and proving the absence of arsenicum in the filtrate by applying some of the foregoing tests.

*Mode of Action of the Antidote.*—The action of the carbonaate of sodium or the magnesia is to precipitate ferric hydrate ($Fe_2 6HO$)—chloride of sodium (NaCl) or magnesium ($MgCl_2$) being formed, which are harmless, if not beneficial, under the circumstances. The reaction between the ferric hydrate and the arsenic results in the formation of insoluble ferrous arseniate.

$$\underset{\text{Ferric hydrate.}}{2(Fe_2 6HO)} + \underset{\text{Arsenic.}}{As_2O_3} = \underset{\text{Ferrous arseniate.}}{Fe_2 2AsO_4} + \underset{\text{Water.}}{5H_2O} + \underset{\text{Ferrous hydrate.}}{Fe_2 2HO}$$

As already stated, dried ferric hydrate (then become an oxyhydrate, $Fe_4O_4 4HO$) (*Ferri Peroxidum Hydratum*, B. P.) has less action on arsenic. Even the moist recently prepared hydrate ($Fe_2 6HO$) loses much of this power as soon as it has become converted into an oxyhydrate ($Fe_4O_3 6HO$), a change which occurs though the hydrate be kept under water (Procter). According to T. and H. Smith this decomposition occurs gradually, but in an increasing ratio: so that after four months the power of the moist mass is reduced to one-half, and after five months to one-fourth.

The so-called Solution of Dialyzed Iron (see Index) is also, as might be expected from its composition, an antidote to arsenic. It should be administered with a little bicarbonate of sodium or bicarbonate of potassium, or any other salt which serves to neutralize the acid that may be present.

---

## QUESTIONS AND EXERCISES.

249. What is the formula of a molecule of arsenicum?
250. In what form does arsenicum occur in nature?
251. Describe the characters of white arsenic.
252. Name the official preparations of arsenicum.
253. What proportion of arsenic ($As_2O_3$) is contained in *Liquor Arsenicalis*, B. P., and *Liquor Arsenici Hydrochloricus*, B. P.?
254. By what method may arsenic be reduced to arsenicum?
255. Give the formulæ of arsenious and arsenic acids and anhydrides.

256. Explain, by diagrams, the reactions which occur in converting arsenic into Arseniate of Sodium by the process of the British Pharmacopœia.

257. Why is anhydrous instead of crystallized arseniate of sodium employed in the preparation of *Liquor Sodæ Arseniatis*, B. P. ?

258. In the preparation of Arseniate of Iron from ferrous sulphate and arseniate of sodium, why is acetate of sodium included ?

259. Describe the manipulations necessary to obtain arsenic in its characteristic crystalline form.

260. How is Reinsch's test for arsenicum applied, and under what circumstances may its indications be fallacious ?

261. Give the details of Marsh's test for arsenicum, and the precautions to be observed in its performance. Explain the reactions by diagrams.

262. What peculiar value has Fleitmann's test for arsenicum ?

263. Describe the conditions under which sulphuretted hydrogen becomes a trustworthy test for arsenicum ?

264. How may a trace of sulphide of arsenicum be detected in sulphur ?

265. How are the salts of copper and silver applied as reagents for the detection of arsenicum ?

266. How are arsenites distinguished from arseniates ?

267. Mention the best antidote in cases of poisoning by arsenic, explain the process by which it may be most quickly prepared, and describe its action ?

268. What light does the action of arsenic on ferric hydrate throw on the constitution of the latter substance ?

---

## ANTIMONY.

Symbol Sb (Stibium). Atomic weight 122.

*Sources and Uses.*—Antimony occurs in nature chiefly as sulphide, $Sb_2S_3$. The *crude* or *black antimony* of pharmacy is this native sulphide freed from earthy impurities by fusion; it has a striated, crystalline, lustrous fracture; subsequently powdered it forms the grayish-black, crystalline *Antimonium nigrum*, B. P., *Antimonii sulphuretum*, U. S. P. The metal is easily obtained from the sulphide by roasting, and then reducing with charcoal and carbonate of sodium. Metallic antimony is an important constituent of *Type-metal*, *Britannia metal* (tea and coffee pots, spoons, etc.), and the best varieties of *Pewter*. The old *pocula emetica*, or everlasting emetic cups, were made of antimony; wine kept in them for a day or two acquired a variable amount of emetic quality. The metal is not used in making the antimonial preparations of the Pharmacopœia, the sulphide alone being, directly or indirectly, employed for this purpose.

Antimony has very close chemical analogies with arsenicum. Its

atom, in the common salts, exerts trivalent activity (*e. g.*, $SbCl_3$), but sometimes it is quinquivalent (*e. g.*, $SbCl_5$).

Antimony, like arsenicum, unites with iodine to form a tri-iodide ($SbI_3$). A bromide ($SbBr_3$) is also known.

## Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

### (*a*) *Reactions having Synthetical Interest.*

### Chloride of Antimony. Antimonious Chloride.

*First Synthetical Reaction.*—Boil half an ounce or less of sulphide of antimony with four or five times its weight of hydrochloric acid in a dish in a fume-chamber or the open air; sulphuretted hydrogen is evolved, and solution of chloride of antimony, $SbCl_3$, obtained.

$$\underset{\text{Sulphide of antimony.}}{Sb_2S_3} + \underset{\text{Hydrochloric acid.}}{6HCl} = \underset{\text{Chloride of antimony.}}{2SbCl_3} + \underset{\text{Sulphuretted hydrogen.}}{3H_2S}$$

This solution, cleared by subsidence, is what is commonly known as *Butter of Antimony* (*Liquor Antimonii Chloridi*, B. P.). If pure sulphide has been used in its preparation, the liquid is nearly colorless; but much of that met with in veterinary pharmacy is simply a by-product in the generation of sulphuretted hydrogen from native sulphide of antimony and hydrochloric acid, and is more or less brown from the presence of chloride of iron. It not unfrequently darkens in color on keeping; this is due to absorption of oxygen from the air and conversion of light-colored ferrous into dark brown ferric chloride or oxychloride.

*True butter of antimony* ($SbCl_3$) is obtained on evaporating the above solution to a low bulk, and distilling the residue. The butter condenses (as a white crystalline semi-transparent mass in the neck of the retort); at the close of the operation it may be easily melted and run down in a bottle, which should be subsequently well-stoppered.

*Pentachloride of antimony* ($SbCl_5$), or *antimonic chloride*, is a fuming liquid, obtained on passing chlorine over the lower chloride.

### Oxychloride of Antimony. Antimonious Oxychloride.

*Second Synthetical Reaction.*—Pour the solution of chloride of antimony produced in the last reaction into several ounces of water; a white precipitate of oxychloride of antimony ($2SbCl_3,5Sb_2O_3$) falls, some chloride of antimony remaining in the supernatant acid liquid.

This is the old *pulvis Algarothi*, *pulvis angelicus*, or *mercurius vitæ*. On standing under water it gradually becomes crystalline.

$$\underset{\text{Chloride of antimony.}}{12SbCl_3} + \underset{\text{Water.}}{15H_2O} = \underset{\text{Oxychloride of antimony.}}{2SbCl_3,5Sb_2O_3} + \underset{\text{Hydrochloric acid.}}{30HCl}$$

## Oxide of Antimony. Antimonious Oxide.

Well wash the precipitate with water, by decantation (*vide* p. 109), and add solution of carbonate of sodium; the chloride remaining with the oxide is thus decomposed, and oxide of antimony ($Sb_2O_3$) alone remains. This is *Antimonii Oxidum* B. P. and U. S. P. It is of a light buff or grayish-white color, or quite white if absolutely free from iron, insoluble in water, soluble in hydrochloric acid, fusible at a low red heat. The moist oxide of antimony may be well washed and employed for the next reaction, or dried over a water-bath. At temperatures above 212° oxygen is absorbed, and other oxides of antimony formed. The presence of the latter is detected on boiling the powder in solution of acid tartrate of potassium, in which oxide of antimony ($Sb_2O_3$) is soluble, but *antimonic anhydride* ($Sb_2O_5$) and the double oxide or so-called *antimonious anhydride* ($Sb_4O_5$) are insoluble.

$$\underset{\text{Oxychloride of antimony.}}{2SbCl_3,5Sb_2O_3} + \underset{\text{Carbonate of sodium.}}{3Na_2CO_3} = \underset{\text{Oxide of antimony.}}{6Sb_2O_3} + \underset{\text{Chloride of sodium.}}{6NaCl} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

*The higher oxide of antimony* ($Sb_2O_5$), termed antimonic oxide or anhydride, corresponding with arsenic anhydride, is obtained on decomposing the pentachloride by water, or on boiling metallic antimony with nitric acid. The variety obtained from the chloride differs in saturating-power from that obtained from the metal, and is termed metantimonic (μετα, *meta*, beyond).

## Tartar Emetic.

*Third Synthetical Reaction.*—Mix the moist oxide of antimony obtained in the previous reaction with about an equal quantity of cream of tartar (6 of the latter to 5 of the dry oxide) and sufficient water to form a paste; set aside for a day to facilitate complete combination; boil the product with water, and filter; the resulting liquid contains the double tartrate of antimony and potassium ($KSbC_4H_4O_7$), potassio-tartrate of antimony, tartrated antimony, or tartar emetic (emetic, from ἐμέω, *emeō*, I vomit; tartar from Τάρταρος, *tartaros*, see Index).

$$\underset{\text{Acid tartrate of potassium.}}{2KHC_4H_4O_6} + \underset{\text{Oxide of antimony.}}{Sb_2O_3} = \underset{\text{Tartar emetic.}}{2KSbC_4H_4O_7} + \underset{\text{Water.}}{H_2O}$$

On evaporation the salt is obtained in colorless transparent triangular-faced crystals of the above composition, with a molecule of water of crystallization, forming the *Antimonium Tartaratum*, B. P. ($KSbC_4H_4O_7,H_2O$), the *Antimonii et Potassii Tartras*, U. S. P.

The formula for tartar emetic is apparently inconsistent with the general formula for tartrates ($R'R'C_4H_4O_6$); this will be subsequently fully explained in connection with Tartaric Acid. The salt appears to be an oxytartrate ($KSbC_4H_4O_6O$).

Tartar emetic is soluble in water, and slightly so in proof-spirit. Dissolved in sherry wine it forms the official *Vinum Antimoniale*, B. P., and *Vinum Antimonii*, U. S. P. It may be externally applied as an ointment, *Unguentum Antimonii Tartarati*, B. P. (*Unguentum Antimonii*, U. S. P.).

### Sulphurated Antimony. Oxysulphide of Antimony.

*Fourth Synthetical Reaction.*—Boil a few grains of sulphide of antimony with solution of soda (potash, U. S. P.) in a test-tube, and filter (or larger quantities in larger vessels, 10 ounces of sulphide to 4½ pints of the official solution of soda, for 2 hours, frequently stirring, and occasionally replacing water lost by evaporation). Into the filtrate, before cool, stir diluted sulphuric acid until the liquid is slightly acid to test-paper; a brownish-red precipitate of oxysulphide of antimony, the *Antimonium Sulphuratum*, B. P. and U. S. P., falls; filter, wash, and dry over a water-bath. It is a mixture of sulphide of antimony ($Sb_2S_3$) with a small and variable amount of oxide ($Sb_2O_3$). The oxide results from the double decomposition of sulphide of antimony and soda.

*Antimonii Oxysulphuretum.*—The United States Pharmacopœia in another preparation orders carbonate of sodium instead of the caustic alkali, and directs that only that precipitate be retained which separates from the alkaline liquid on cooling.

If a small quantity of sulphur be boiled with the sulphide of antimony in solution of soda, the precipitate on addition of sulphuric acid will have a much brighter color, chiefly on account of the presence of a higher sulphide having a yellow color ($Sb_2S_5$).

These are some of the many varieties of *mineral kermes*, so called from their similarity in color to the *insect kermes*. Kermes is the name, now obsolete, of the *Coccus Ilicis*, a sort of cochineal-insect, full of reddish juice, and used for dyeing from the earliest times. The color of the precipitate is affected by the temperature as well as state of dilution of the alkaline liquid when the acid is added. When

the alkaline liquid is boiled, especially if long exposed to air, oxygen is absorbed by some of the antimony, whose sulphur uniting with the trisulphate forms a portion of the lighter yellow pentasulphide. *Kermes mineral* thus varies much in proportion of oxide and of pentasulphide as well as in the physical condition of its trisulphide.

*Explanation of Process.*—The sulphides and oxides of antimony, like those of arsenicum, react with the sulphides and oxides of certain metals to form soluble salts ($Na_3SbS_3$ and $Na_3SbO_3$). The former is deposited in yellow tetrahedral crystals when such an alkaline solution is set aside to slowly cool.

$$\underset{\text{Sulphide of antimony.}}{2Sb_2S_3} + \underset{\text{Soda.}}{6NaHO} = \underset{\text{Sulph-antimonite of sodium.}}{2Na_3SbS_3} + \underset{\text{Oxide of antimony.}}{Sb_2O_3} + \underset{\text{Water.}}{3H_2O}$$

$$\underset{\text{Oxide of antimony.}}{Sb_2O_3} + \underset{\text{Soda.}}{6NaHO} = \underset{\text{Antimonite of sodium.}}{2Na_3SbO_3} + \underset{\text{Water.}}{3H_2O}$$

In the hot solutions of these salts sulphide and oxide of antimony are soluble, and are reprecipitated in an indefinite state of combination, partially, on cooling, or wholly on the addition of acid. The acid also decomposes the oxysalt with precipitation of oxide, and the sulphur salt with precipitation of orange sulphide of antimony. The acid is added to the liquid before much oxysulphide has deposited (that is, before the solution is cool), in order to insure uniformity of product.

$$\underset{\text{Sulph-antimonite of sodium.}}{2Na_3SbS_3} + \underset{\text{Sulphuric acid.}}{3H_2SO_4} = \underset{\text{Sulphate of sodium.}}{3Na_2SO_4} + \underset{\text{Sulphide of antimony.}}{Sb_2S_3} + \underset{\text{Sulphuretted hydrogen.}}{3H_2S}$$

$$\underset{\text{Antimonite of sodium.}}{2Na_3SbO_3} + \underset{\text{Sulphuric acid.}}{3H_2SO_4} = \underset{\text{Sulphate of sodium.}}{3Na_2SO_4} + \underset{\text{Oxide of antimony.}}{Sb_2O_3} + \underset{\text{Water.}}{3H_2O}$$

The oxide and sulphide indicated in these equations, together with excess of sulphide of antimony originally dissolved by the alkaline liquid, are all precipitated when the acid is added, and form the *Sulphurated Antimony* of the Pharmacopœia, "an orange-red powder, readily dissolved by caustic soda, also by hydrochloric acid with the evolution of sulphuretted hydrogen and the separation of a little sulphur." Its antimony is detected by dissolving the powder in hydrochloric acid or in solution of acid tartrate of potassium, and passing sulphuretted hydrogen through the liquid, as described in the first analytical reaction.

---

The previous four synthetical reactions illustrate the official processes for the respective substances. The solution of chloride of antimony is only used in the preparation of oxide; the oxide, besides its use in the preparation of tartar emetic, is mixed with twice its weight of phosphate of calcium (purified bone-earth) to form *Pulvis Antimonialis*, B. P.

The *sulphides* and *hydride* of antimony are incidentally mentioned in the following analytical paragraphs.

### (*b*) *Reaction having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—Through an acidified antimonial solution pass sulphuretted hydrogen; an orange precipitate of amorphous sulphide of antimony falls. It has the same composition as the crystalline black sulphide ($Sb_2S_3$), into which, indeed, when dried, it is quickly converted by heat. Like sulphide of arsenicum, it is soluble in alkaline solutions. Collect a portion on a filter, and, when well drained, add *strong* hydrochloric acid; it dissolves—unlike sulphide of arsenicum.

A higher sulphide of antimony ($Sb_2S_5$), corresponding to the higher sulphide of arsenicum, exists. It is formed on passing sulphuretted hydrogen through an acidified solution of the higher chloride ($SbCl_5$), or on boiling black sulphide of antimony and sulphur with an alkali, and decomposing the resulting filtered liquid by an acid.

*Note.*—The arsenious and antimonious compounds are those chiefly employed in medicine; arseniates of sodium and iron are, however, sometimes employed. The arseniates and rarely an antimoniate are useful in analysis, and the antimonic chloride in chemical research. The higher compounds of both elements are noticed here chiefly to draw attention to the close analogy existing between arsenicum and antimony, an analogy carried out in the numerous other compounds of these elements.

*Second Analytical Reaction.*—Dilute two or three drops of the solution of chloride of antimony with water; a precipitate of oxychloride occurs, the formation of which has been explained under the similar synthetical reaction. The occurrence of this precipitate distinguishes antimony from arsenicum, but is a reaction that cannot be fully relied upon in analysis, because requiring the presence of too much material and the observance of too many conditions. Add a sufficient quantity of hydrochloric acid to dissolve the precipitate, and boil a piece of copper in the solution as directed in the corresponding test for arsenicum (*vide* page 172); antimony is deposited on the copper. Wash, dry, and heat the copper in a test-tube as before; the antimony, like the arsenicum, is volatilized off the copper and condenses on the side of the tube as white oxide, but the sublimate, from its low degree of volatility, condenses close to the copper, moreover, it is destitute of crystalline character—that is to say, it is *amorphous* (α, *a*, without; μορφή, *morphē*, shape).

Shake out the copper, and boil water in the tube for several minutes. Do the same with the arsenical sublimate similarly obtained. The deposit of arsenic slowly dissolves, and may be

recognized in the solution by ammonio-nitrate of silver; the antimonial sublimate is insoluble.

*Third Analytical Reaction.*—Perform the experiments described under Marsh's test for arsenicum (pp. 173, 174), carefully observing all the details there mentioned, but using a few drops of solution of chloride of antimony or tartar emetic instead of the arsenical solution. Antimoniuretted hydrogen, or hydride of antimony ($SbH_3$), is formed and decomposed in the same way as arseniuretted hydrogen.

To one of the arsenicum spots on the porcelain lid (p. 174) add a drop of a solution of "chloride of lime" (bleaching-powder); it quickly dissolves. Do the same with an antimony spot; it is unaffected.

Heat more quickly causes the volatilization of an arsenicum than an antimony spot; sulphydrate of ammonium more readily dissolves the antimony than the arsenicum.

Boil water for several minutes in the beaker or wide test-tube containing the arsenious sublimate (page 174); it slowly dissolves, and may be recognized in the solution by the yellow precipitate given on the addition of solution of ammonio-nitrate of silver. The antimonial sublimate, similarly treated, gives no corresponding reaction.

Pass a slow current of sulphuretted hydrogen through the delivery-tube removed from the hydrogen-apparatus (page 174), and, when the air may be considered to have been expelled from the tube, gently heat that portion containing the deposit of arsenicum; the latter will be converted into a *yellow* sublimate of sulphide of arsenicum. Remove the tube from the sulphuretted-hydrogen apparatus, and repeat the experiment with a similar antimony deposit; it is converted into *orange* sulphide of antimony, which, moreover, owing to inferior volatility, condenses nearer to the flame than sulphide of arsenicum.

Pass dry hydrochloric acid gas through the two delivery-tubes. This is accomplished by adapting first one tube and then the other by a cork to the test-tube containing a few lumps of common salt, on which a little sulphuric acid is poured during the momentary removal of the cork. The sulphide of antimony dissolves and disappears; the sulphide of arsenicum is unaffected.

*Thorough perception of the chemistry of arsenicum and antimon will be obtained on constructing equations or diagrams descriptiv of each of the foregoing reactions.*

*Antidote.*—The introduction of poisonous doses of antimonials into the stomach is fortunately quickly followed by vomiting.

If vomiting has not occurred, or apparently to an insufficient extent, any form of tannic acid may be administered (infusion of tea, nutgalls, cinchona, oak-bark, or other astringent solutions or tinctures), an insoluble tannate of antimony being formed, and absorption of the poison consequently somewhat retarded. The stomach-pump must be applied as quickly as possible.

Recently precipitated moist ferric hydrate is also, according to T. and H. Smith, a perfect absorbent of antimony from its solutions, the chemical actions being probably, they say, similar to that which takes place between ferric hydrate and ansenious anhydride. It may be given in the form of a mixture of perchloride of iron with either carbonate of sodium or magnesia.

These statements may be verified by mixing together the various substances, filtering, and testing the filtrate for antimony in the usual manner.

---

DIRECTIONS FOR APPLYING THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE OF THE ELEMENTS ARSENICUM AND ANTIMONY.

Acidify the liquid with hydrochloric acid, and pass through it sulphuretted hydrogen :—

A *yellow* precipitate indicates arsenicum.

An *orange* precipitate indicates antimony.

The result may be confirmed by the application of other tests.

---

DIRECTIONS FOR APPLYING THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF BOTH ARSENICUM AND ANTIMONY.

Acidify a small portion of the liquid with hydrochloric acid, and pass through it sulphuretted hydrogen.

*Note* I. If the precipitate by sulphuretted hydrogen is unmistakably orange, antimony may be put down as present, and arsenicum only further sought by the application of Fleitmann's test to the solution of the sulphide in aqua regia* freed from sulphur by boiling, or, better, to the original solution.

*Note* II. Sulphide of antimony is far less readily soluble than sulphide of arsenicum in solution of carbonate of ammonium. But this

---

* *Aqua regia* is a mixture of four parts hydrochloric and three parts nitric acid. It was so called, from its property of dissolving gold, the "king" of metals.

fact possesses limited analytical value; for the color of the sulphides is already sufficient to distinguish the one from the other when they are unmixed; and when mixed, much sulphide of antimony will prevent a little sulphide of arsenicum from being dissolved by the alkaline carbonate, while much sulphide of arsenicum will carry a little sulphide of antimony into the solution. When the proportions are, apparently, from the color of the precipitate, less wide, solution of carbonate of ammonium will be found useful in roughly separating the one sulphide from the other. On filtering and neutralizing the alkaline solution by an acid, the yellow sulphide of arsenicum is reprecipitated. The orange sulphide of antimony will remain on the filter.

*Note* III. Solution of bisulphite of potassium is said by Wöhler to be a good reagent for separating the sulphides of arsenicum and antimony, the former being soluble, the latter insoluble in the liquid.

*Note* IV. Another reagent for separating the sulphides of arsenicum and antimony is *strong* hydrochloric acid. As little water as possible must be present. On boiling, the sulphide of antimony dissolves, while the sulphide of arsenicum remains insoluble. The liquid slightly diluted, filtered, more water added, and sulphuretted hydrogen again transmitted, gives orange sulphide of antimony. The process should previously be tried on the precipitated mixed sulphides. The presence of arsenicum may be confirmed by the application of Fleitmann's test to the original solution.

*Note* V. If the precipitate by sulphuretted hydrogen is unmistakably yellow, arsenicum may be put down as present, and any antimony detected by the previous or one of the following two processes. These two processes are rather long, and require much care in their performance, but are useful, because a small quantity of antimony in much arsenicum, or *vice versâ*, may be detected by their means.

*First Process.*—Generate hydrogen and pass it through a small wash-bottle containing solution of acetate of lead, to free the gas from any trace of sulphuretted hydrogen it may possess, and then through a dilute solution of nitrate of silver contained in a test-tube. When the apparatus is in good working order, pour into the generating-bottle the solution to be examined, adding it gradually to prevent violent action. After the gas has been passing for five or ten minutes, examine the contents of the nitrate-of-silver tube; arsenicum, if present, will be found in the solution in the state of arsenious acid,

$$AsH_3 + 3H_2O + 6AgNO_3 = H_3AsO_3 + 6HNO_3 + 3Ag_2;$$

while antimony, if present, will be found in the black precipitate that has fallen, according to the following equation:—

$$SbH_3 + 3AgNO_3 = SbAg_3 + 3HNO_3.$$

The arsenious radical may be detected in the clear, filtered, supernatant liquid, which still contains much nitrate of silver,

by cautiously neutralizing with a very dilute solution of ammonia, or by adding a few drops of solution of ammonio-nitrate of silver, yellow arsenite of silver being produced. The antimony may be detected by washing the black precipitate, boiling it in an open dish with solution of tartaric acid, filtering, acidulating with hydrochloric acid, and passing sulphuretted hydrogen through the solution—the orange sulphide of antimony being precipitated (Hofmann).

*Second Process.*—Obtain the metallic deposit in the middle of the delivery-tube as already described under Marsh's test. Act on the deposit by sulphuretted hydrogen gas, and then by hydrochloric acid gas as detailed in the third analytical reaction of antimony (p. 185). If both arsenicum and antimony are present, the deposit, after the action of sulphuretted hydrogen, will be found to be of two colors, the yellow sulphide of arsenicum being usually further removed from the heated portion of the tube than the orange sulphide of antimony. Moreover, subsequent action of hydrochloric acid gas causes disappearance of the antimonial deposit, which is converted into chloride of antimony and carried off in the stream of gas.

The chief objection to this process is the liability of the operator mistaking sulphur, deposited from the sulphuretted hydrogen gas by heat, for sulphide of arsenicum. But the presence or absence of arsenicum is easily confirmed by applying Fleitmann's test to the original solution, while the process is most useful for the detection of a small quantity of salt of antimony when mixed with much arsenical compounds.

---

The laboratory student may now proceed to the analysis of aqueous solution of salts of any of the metallic elements hitherto considered. The method followed may be that for the separation of the previous three groups, sulphuretted hydrogen being first passed through the solution to throw out arsenicum and antimony. The whole scheme of analysis is given on the next page. Three or four solutions should be examined before proceeding to the last group of metals.

Learners who have no opportunity of working at practical analysis will gain much knowledge by endeavoring not to remember, but to understand these methods of separating elements from each other in a solution containing several compounds.

TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF **ANY OR ALL** OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.

Acidify with HCl, and pass $H_2S$ through the solution; filter.

<table>
<tr><td rowspan="5">Precipitate<br>As Sb.<br>Wash, separate by carb. ammon.; or by strong HCl; or dissolve in a few drops HCl and $HNO_3$, and examine as described on pages 186 and 187.</td><td colspan="5">Filtrate<br>Fe Al Zn Ba Ca Mg K Na Am.<br>Add AmHO;* AmHS; stir, filter.</td></tr>
<tr><td colspan="2">Precipitate†<br>Fe Al Zn.<br>Wash, dissolve in HCl, boil with $HNO_3$ (p. 153), add KHO in excess,‡ stir, filter.</td><td colspan="3">Filtrate<br>Ba Ca Mg K Na Am.<br>Add $Am_2CO_3$, boil, filter.</td></tr>
<tr><td rowspan="2">Precipitate<br>Fe.<br>Examine orig. solution for ferrous or feric state.</td><td rowspan="2">Filtrate<br>Al Zn.<br>Neut. by HCl. add AmHO, stir, filter.</td><td colspan="2">Precipitate<br>Ba Ca.<br>Dissolve in $HC_2H_3O_2$, add $K_2CrO_4$, filter.</td><td colspan="2">Filtrate<br>Mg K Na Am.<br>Add $Am_2HPO_4$, shake, filter.</td></tr>
<tr><td>Ppt.<br>Al.</td><td>Sol.<br>Zn.</td><td>Precipitate<br>Ba.</td><td>Filtrate<br>Ca.</td><td>Precipitate<br>Mg.</td><td>Filtrate<br>Am Na K.<br>(page 100).</td></tr>
</table>

* It is not usually necessary here to add $NH_4Cl$ to keep Mg in solution, the HCl with some of the $NH_4HO$ commonly giving sufficient $NH_4Cl$ for this purpose.

† Much time may sometimes be saved by carefully remembering the color of the various hydrates and sulphides precipitated. Thus, if the AmHS precipitate is white, iron cannot be present, and AmHO, for Al and Zn, may be at once added to the hydrochloric solution of the precipitate. The group-tests in this Table are $H_2S$, AmHS, and $Am_2CO_3$.

‡ The pupil should vary the process sometimes, as described in the foot-notes on pages 162 and 164. The treatment of other groups should occasionally be varied. By remembering the reactions on which analytical processes are based a pupil may often devise modified processes, and thus frequently save time, avoid working in one groove only, and acquire broader chemical views.

## QUESTIONS AND EXERCISES.

269. What is the composition and source of the *Black Antimony* of pharmacy?

270. In what alloys is metallic antimony a characteristic ingredient?

271. What is the quantivalence of antimony as far as indicated by the formulæ of the official preparations?

272. By a diagram show how "Butter of antimony" is prepared.

273. Write out equations or diagrams expressive of the reactions which occur in converting chloride of antimony into oxide.

274. What is the formula of Tartar Emetic?

275. Explain the official process for the preparation of Oxysulphide of Antimony (*Antimonium Sulphuratum*, B. P. or U. S. P.) by aid of diagrams.

276. Give a comparative statement of the tests for arsenicum and antimony.

277. How is antimony detected in the presence of arsenicum?

278. How may arsenicum and iron be distinguished analytically?

279. Describe a method by which antimony, magnesium, and iron may be separated from each other.

280. Draw out an analytical chart for the examination of an aqueous liquid containing salts of arsenicum, zinc, calcium, and ammonium.

---

## COPPER, MERCURY, LEAD, SILVER.

These metals, like arsenicum and antimony, are precipitated from acidified solutions by sulphuretted hydrogen, in the form of sulphides; but the sulphides, unlike those of arsenicum and antimony, are insoluble in alkalies. The atom of copper is usually bivalent, Cu″; mercury bivalent in the mercuric salts, Hg″, and univalent in the mercurous salts, Hg′; lead sometimes quadrivalent, Pb⁗, but generally exerting only bivalent activity, Pb″; and silver univalent, Ag′.

### COPPER.

Symbol Cu. Atomic weight 63.5.

*Source.*—The commonest ore of this metal is *copper pyrites*, a double sulphide of copper and iron, raised in Cornwall; Australia and Russia supply *malachite*, a mixed carbonate and hydrate; much ore is also imported from South America. It is smelted in enormous quantities at Swansea, South Wales, a locality peculiarly fitted for the operation on account of its proximity to the coal-fields, and its position as a sea-coast town—these advantages at all times insuring cheap fuel and freightage to the different metallurgical establishments.

*Alchemy.*—The alchemists termed this metal *Venus*, perhaps on account of the beauty of its lustre, and gave it her symbol ♀, a compound hieroglyphic also indicating a mixture of gold ☉ and a certain hypothetical substance called acrimony ✠, the corrosive nature of which was symbolized by the points of a Maltese cross. To this day the blue show-bottle in the shop-window of the pharmacist is occasionally ornamented by such a symbol, indicative, possibly, of the fact that the blue liquid in the vessel is a preparation of copper.

*Coinage.*—The material of British copper coinage is now a bronze mixture composed in 100 parts by weight of 95 copper, 4 tin, and 1 zinc, the same as in the copper coinage of France. The penny is coined at the rate of 48 pence in one pound avoirdupois, of 7000 grains, or 453.6 grammes; the halfpenny at 80 in the pound avoirdupois, and the farthing at 160. British bronze coins are a legal tender in payments to the amount of 1*s.*

*Metallic Copper* (*Cuprum*, B. P. and U. S. P.) in the form of fine wire, about No. 25, is used in prepararing *Spiritus Ætheris Nitrosi*, B. P. and U. S. P. *Copper Foil*, B. P., is "pure metallic copper, thin and bright."

*Quantivalence.*—Copper forms two classes of salts; in one the atom is bivalent ($Cu''$), in the other exerts univalent activity ($Cu_2''$). The former are of primary importance, the latter being for the most part unstable and wanting in technical interest. Their compounds are distinguished as cupric and cuprous; but those of the higher class only have general interest, and will be almost exclusively alluded to in the following paragraphs. Cuprous iodide ($Cu_2I_2$) will subsequently be referred to as a convenient form in which to remove iodine from solution, and the formation of cuprous oxide ($Cu_2O$), under given circumstances, as an indicator of the presence of sugar in a liquid.

## Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

### (*a*) *Synthetical Reactions.*

The processes for the following salts include the only synthetical reactions having any medical or pharmaceutical interest: 1, cupric oxide, the black oxide of copper, prepared by heating a piece of copper to low redness on a piece of earthenware in an open fire; 2, cupric sulphate, the common sulphate of copper, prepared by boiling black oxide and about an equal weight of sulphuric acid in water, filtering, and setting aside the solution so that crystals may form on cooling; and 3, ammonio-sulphate of copper, for the preparation of which see p. 177; also p. 193.

$$\underset{\text{Copper.}}{Cu_2} + \underset{\text{Oxygen.}}{O_2} = \underset{\substack{\text{Cupric}\\\text{oxide.}}}{2CuO}$$

$$\underset{\substack{\text{Cupric}\\\text{oxide.}}}{CuO} + \underset{\substack{\text{Sulphuric}\\\text{acid.}}}{H_2SO_4} = \underset{\substack{\text{Cupric}\\\text{sulphate.}}}{CuSO_4} + \underset{\text{Water.}}{H_2O}$$

*Sulphate of Copper* (*Cupri Sulphas*, B. P. and U. S. P.) ($CuSO_4$, $5H_2O$), *blue vitriol*, *bluestone*, or *cupric sulphate*, is the only copper salt of much importance in Pharmacy. It is a by-product in silver-refining ($2Ag_2SO_4 + Cu_2 = 2CuSO_4 + 2Ag_2$). It is also formed by roasting copper pyrites. In the latter operation sulphide of iron and sulphide of copper are oxidized to sulphates; but the low red heat finally employed decomposes the sulphate of iron, while the sulphate of copper is unaffected; it is purified by crystallization from a hot aqueous solution, though frequently much sulphate of iron remains in the crystals. Sulphate of copper results on dissolving in diluted sulphuric acid the black oxide (CuO) obtained in annealing copper plates (see above equations); it may also be prepared by boiling copper with three times its weight of sulphuric acid ($2H_2SO_4 + Cu = CuSO_4 + SO_2 + 2H_2O$), diluting, filtering, evaporating, and crystallizing. In this process a little black sulphide of copper is formed.

*Anhydrous Sulphate of Copper* ($CuSO_4$) is a yellowish-white powder prepared by depriving the ordinary blue crystals of sulphate of copper of their water of crystallization by exposing them to a temperature of about 400° F. It is used in testing alcohol and similar spirituous liquids for water, becoming blue if the latter be present.

*Verdigris* (from *verde-gris*, Sp. green-gray) is a Subacetate (*Cupri Subacetas*, U. S. P.) or Oxyacetate of Copper (B. P.) ($Cu_2O$-$2C_2H_3O_2$), obtained by exposing alternate layers of copper and fermenting refuse grape-husks to the action of air. Digested with twice its weight of acetic acid and a little water, the mixture being evaporated to dryness and the residue dissolved in water, it forms the official Solution of Acetate of Copper ($Cu2C_2H_3O_2$).

The modes of forming *cupric sulphide*, *hydrate*, *oxide*, *ferrocyanide*, and *arsenite*, as well as the precipitation of *metallic copper*, are incidentally alluded to in the following analytical paragraphs.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.* — Pass sulphuretted hydrogen through an acidified solution of a copper salt (sulphate, for example); black cupric sulphide (CuS) falls.

*Second Analytical Reaction.* — Add sulphydrate of ammonium to an aqueous copper solution; cupric sulphide is again precipitated, insoluble in excess.

*Note.* — Cupric sulphide is not altogether insoluble in sulphydrate of ammonium if free ammonia or much ammoniated salt be present; it is quite insoluble in the fixed alkaline sulphides.

*Third Analytical Reaction.* — Immerse a piece of iron or steel, such as the point of a penknife or a piece of wire, in a few drops of a copper solution; the copper is deposited, of characteristic color, an equivalent quantity of iron passing into solution.

By this reaction copper may be recovered on the large scale from waste solutions, old hoop or other scrap iron being thrown into the liquors.

*Fourth Analytical Reaction.*—Add ammonia to a cupric solution; cupric hydrate (Cu2HO) of a light-blue color is precipitated. Add excess of ammonia; the precipitate is redissolved, forming a blue solution of ammonio-salt of copper, so deep in color as to render ammonia an exceedingly delicate test for this metal.

An ammonio-sulphate of copper may be obtained in large crystals by adding strongest solution of ammonia to powdered sulphate of copper until the salt is dissolved, placing the liquid in a test-glass or cylinder, cautiously pouring in twice its volume of strong alcohol or methylated spirit, taking care that the liquids do not become mixed, tying over the vessel with bladder, and setting aside for some weeks is a cool place. (Wittstein.) The constitution of ammonio-sulphate and other ammonio-salts of copper and corresponding salts of silver will be alluded to in connection with "white precipitate," the official "ammoniated mercury."

*Cuprum Ammoniatum*, U. S. P., is an ammonio-sulphate of copper prepared by rubbing together sulphate of copper and carbonate of ammonium until effervescence ceases, and drying the product.

*Fifth Analytical Reaction.*—Add solution of potash or soda to a cupric solution; cupric hydrate (Cu2HO) is precipitated, insoluble in excess. Boil the mixture in the test-tube; the hydrate is decomposed, losing the elements of water, and becoming the black anhydrous oxide (CuO).

*Sixth Analytical Reaction.*—Add solution of ferrocyanide of potassium ($K_4Fcy$) to an aqueous cupric solution; a reddish-brown precipitate of cupric ferrocyanide ($Cu_2Fcy$) falls. This also is a delicate test for copper.

*Seventh Analytical Reaction.*—To a cupric solution add solution of arsenic, and cautiously neutralize with alkali; green cupric arsenite ($CuHAsO_3$) falls.

*Note.*—This precipitate has been already mentioned under arsenicum. An arsenicum salt is thus a test for copper as a copper salt is for arsenicum—a remark that may obviously be extended to most analytical reactions; for the *body acted upon characteristically by a reagent is as good a test for the reagent as the reagent is for it;* indeed it becomes a reagent when the other body is the object of search.

*Antidotes.*—In cases of poisoning by compounds of copper, iron filings should be administered, the action of which has just been explained (see third analytical reaction). Ferrocyanide of potassium may also be given (see sixth analytical reaction).

Albumen forms with copper a compound insoluble in water; hence raw eggs should be swallowed, vomiting being induced or the stomach-pump applied as speedily as possible.

## QUESTIONS AND EXERCISES.

281. What are the relations of copper, mercury, lead, and silver to each other and to arsenicum and antimony?
282. Name the sources of copper.
283. What proportion of copper is contained in English and French "copper" coins?
284. Give diagrams showing how Sulphate of Copper is prepared on the small and large scales.
285. Work out a sum showing how much Crystallized Sulphate of Copper may be obtained from 100 parts of sulphide.—*Ans.* 261¼.
286. How may Oxide of Copper be prepared?
287. Mention the formula of Verdigris.
288. Name a good clinical test for copper.
289. What is the analytical position of copper?
290. Mention the chief tests for copper.
291. How may copper be separated from arsenicum?
292. Why is finely divided iron an effective antidote in cases of poisoning by Copper?

## MERCURY.

Symbol Hg. Atomic weight 200.

Molecular weight 200 (*not* double the atomic weight).

*Source.*—Mercury occurs in nature as sulphide (HgS), forming the ore *cinnabar* (an Indian name expressive of something red), and is obtained from Spain, California, Eastern Hungary, China, Japan, and Peru.

*Preparation.*—The metal is separated by roasting off the sulphur and then distilling, or distilling with lime, which combines with and retains the sulphur.

*Properties.*—Mercury (*Hydrargyrum*, B. P. and U. S. P.) is a silver-white lustrous metal, liquid at common temperatures. It boils at 662° F., and at —40° F., solidifies to a malleable mass of octahedral crystals. When quite free from other metals it does not tarnish, and its globules roll freely over a sheet of white paper without leaving any streak or losing their spherical form.

*Formula.*—The formula of the mercury molecule is Hg and not $Hg_2$, because (at all events at the high temperature at which alone the weight of its vapor can be determined) two volumes, which if hydrogen would weigh two parts ($H_2$) or oxygen thirty-two parts ($O_2$), in the case of mercury vapor weigh only two hundred parts

(Hg); that is, only once the atomic weight, not twice. That 200, and not 100, is the atomic weight of mercury is shown by the fact that 200 is the minimum proportion relative to 1 of hydrogen in which mercury combines, and by its relations to heat. Still it is difficult to imagine an atom existing in the free state in nature; and the suggestion has been made that (as is proved to be the case with sulphur) mercury, as we know it, is in abnormal condition, and that if the weight of its vapor could be taken at a lower temperature, or under some other condition, its molecular weight might be found to be 400. Similar remarks may be made respecting zinc, the molecular weight of which, so far as we know, is identical with its atomic weight.

*Medicinal Compounds.*—The compounds of mercury used in medicine are all obtained from the metal. The metal itself, rubbed with chalk or with confection of roses and powdered liquorice-root, or with lard and suet, until globules are not visible *to the unaided eye*, is often used in medicine. The preparations are: the *Hydrargyrum cum Creta*, B. P. and U. S. P., or "Gray Powder;" *Pilula Hydrargyri*, B. P. and U. S. P., or "Blue Pill;" and *Unguentum Hydrargyri* B. P. and U. S. P., or "Blue Ointment." There are also a Compound Ointment, a Plaster of Mercury, a Plaster of Ammoniacum and Mercury, a Liniment, and a Suppository. Their therapeutic effects are probably due not to the large quantity of metallic mercury in them, but to the small quantities of black and red oxide which occur in them through the action of the oxygen of the air on the finely-divided metal. The proportion of oxide or oxides varies according to the age of the specimen.

All these medicinal preparations of metallic mercury are indefinite and unsatisfactory, and that through no fault of the pharmacist. They much need investigation by therapeutists. Here, as in many similar cases, if medicine would first ascertain her own requirements and then make them known, her handmaid pharmacy would be found quite capable of supplying them.

*Mercurous and Mercuric Compounds.*—Mercury combines with other elements and radicals in two proportions; those compounds in which the other, acidulous, radicals are in the lesser amount are termed *mercurous*, the higher being *mercuric*. Thus, calomel ($HgCl$*) is mercurous chloride, while corrosive sublimate ($HgCl_2$) is mercuric chloride. In every pair of mercuric compounds the mercuric contains *twice* as much complementary radical, in proportion to the mercury, as the mercurous.

*Note on Nomenclature.*—The remarks made concerning the two classes of iron salts, ferrous and ferric (p. 141), apply in the main to the two series of mercury salts. The latter are systematically distinguished in most modern works by the terms *mercurous* and *mercuric*. In the British and United States Pharmacopœias, however,

---

* The specific gravity of the vapor of calomel, and the fact that the salt is *not* decomposed at the temperature at which its specific gravity is taken, indicate that the formula of calomel is $HgCl$, and not $Hg_2Cl_2$.

which include only a few in comparison with the whole number of mercury salts, older and more strongly contrasted names are employed, thus:—

| Systematic names. | Official names. |
|---|---|
| Mercurous iodide . . . . | Green iodide of mercury. |
| Mercuric iodide . . . . . | Red iodide of mercury. |
| Mercurous nitrate . . . . | Not mentioned in B. P. |
| Mercuric nitrate . . . . . | Nitrate of mercury. |
| Mercurous sulphate . . . . | Not mentioned in B. P. |
| Mercuric sulphate . . . . | Sulphate of mercury. |
| Mercurous chloride . . . . | Subchloride of mercury. |
| Mercuric chloride . . . . | Perchloride of mercury. |
| Mercurous oxide . . . . | Black oxide of mercury. |
| Mercuric oxide . . . . . | Red oxide of mercury. |

*Specific Gravity.*—Mercury is 13.6 times as heavy as water.

*Amalgams.*—The compound formed in fusing metals together is usually termed an *alloy* (*ad* and *ligo*, to bind); but if mercury is a constituent, an *amalgam* (*μάλαγμα*, *malagma*, from *μαλάσσω*, *malassō*, to soften, the presence of mercury lowering the melting point of such a mixture). Most metals, even hydrogen, according to Leow, form amalgams.

## Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

### (*a*) *Synthetical Reactions.*

### The Two Iodides.

*First Synthetical Reaction.*—Rub together a small quantity of mercury and iodine, controlling the rapidity of combination by adding a few drops of spirit of wine, which, by evaporation, absorbs heat, and thus keeps down temperature. The product is either mercuric iodide, mercurous iodide, or a mixture of the two, as well as mercury or iodine if excess of either has been employed. If the two elements have been previously weighed in single atomic proportions, 200 of mercury to 127 of iodine (about 8 to 5, or 1 ounce of mercury to 278 grains of iodine), the mercurous or green (grayish-green) iodide results (HgI) (*Hydrargyyri Iodidum Viride*, B. P. and U. S. P.); if in the proportion of one atom of mercury to two atoms of iodine (200 to twice 127, or about 4 to 5), the mercuric or red iodide, $HgI_2$, results, an iodide that is also official, but made in another way. (See page 198.) The green iodide should be made and dried (without heat) with as little exposure to light as possible.

Mercurous iodide is decomposed slowly by light, and quickly by heat, into mercuric iodide and mercury. Mercuric iodide occurring as an impurity in mercurous iodide may be detected by digesting in ether (in which mercurous iodide is insoluble), filtering and evaporating to dryness; mercuric iodide remains. Mercuric iodide is stable, and may be sublimed in scarlet crystals without decomposition. (For details of the method by which a specimen of the crystals may be obtained, and the precautions to be observed, *vide* "Corrosive sublimate," p. 200.)

*Relation of Mercuric Iodide to Light.*—In condensing, mercuric iodide is at first yellow, afterwards acquiring its characteristic scarlet color. This may be shown by smearing or rubbing a sheet of white paper with the red iodide, and then holding the sheet before a fire or over a flame for a few seconds. As soon as the paper becomes hot the red instantly changes to yellow, and the salt does not quickly regain its red color, even when cold, if the paper is carefully handled. But if a mark be made across the sheet by anything at hand, or the salt be pressed or rubbed in any way, the portions touched immediately return to the scarlet condition. According to Warrington, this change is consequent upon rhomboidal crystals being converted into octahedra with a square base, and will serve as an excellent illustration of the influence of physical structure in causing color. The yellow modification so acts on the rays of white light shining on its particles as to absorb the violet and reflect the complementary hue, the yellow, which, entering the eye of the observer, strikes his retina, and thus conveys to the brain the impression of yellowness; and the red modification, though actually the same chemical substance, is sufficiently different in the structure of its particles to absorb the green constituent of white light and reflect the complementary ray, the red.

*Illustration of the Chemical Law of Multiple Proportions* (p. 48).—Applying the atomic theory to the above iodides, it will at once be apparent why mercury and iodine should combine in the proportion of 200 of mercury with either 127 or 254 of iodine, and not with any intermediate quantity. For it is part of that theory that masses are composed of atoms, and that atoms are indivisible; and that the weight of the atom of mercury is to that of iodine as 200 is to 127. Mercury and iodine can only combine, therefore, in atomic proportions, atom to atom (which is the same as 200 to 127), or one atom to two atoms (which is the same as 200 to 254). To attempt to combine them in any intermediate proportion would be useless, a mere mixture of the two iodides would result. A higher proportion of mercury than 200 to 127 of iodine gives but a mixture of mercurous iodide and mercury; a higher proportion of iodine than 254 to 200 of mercury gives but a mixture of mercuric iodide and iodine. Or, for example, 200 grains of mercury, mixed with, say, 200 of iodine, would yield 139 grains of mercurous iodide, and 261 grains of mercuric iodide; for the 200 grains of mercury uniting with 127 grains of the iodine give, for the moment, 327 grains of mercurous iodide and 73 grains of iodine still free. The 73 grains of iodine will immediately unite with 188 grains of the mercurous iodide (for if 127

of I require 327 of HgI to form $HgI_2$, 73 will require 188), and form 261 grains of mercuric iodide, diminishing the 327 grains of mercurous iodide to 139 grains.

*Preparation of Red Iodide of Mercury by precipitation.*—To a few drops of a solution of a mercuric salt (corrosive sublimate, for example), add solution of iodide of potassium, drop by drop; a precipitate of mercuric iodide, $HgI_2$, forms, and at first quickly redissolves, but is permanent when sufficient iodide of potassium has been added. Continue the addition of iodide of potassium; the precipitate is once more redissolved.

$$\underset{\text{Mercuric chloride.}}{HgCl_2} + \underset{\text{Iodide of potassium.}}{2KI} = \underset{\text{Mercuric iodide.}}{HgI_2} + \underset{\text{Chloride of potassium.}}{2KCl}$$

*Note.*—When first precipitated, mercuric iodide is yellowish-red, but soon changes to a beautiful scarlet. Its solubility either in solution of the mercuric salt or in solution of iodide of potassium renders the detection of a small quantity of a mercuric salt by iodide of potassium, or a small quantity of an iodide by a mercuric solution, difficult, and hence lessens the value of the reaction as a test. But the reaction has synthetical interest, the method by precipitation being that adopted in the British and United States Pharmacopœias (*Hydrargyri Iodidum Rubrum*, B. P. and U. S. P.). Mercuric iodide thus made has the same composition as that prepared by direct combination of its elements. Equivalent proportions of the two salts must be used in making the preparation ($HgCl_2 = 271$; $2KI = 332$). About 4 parts of corrosive sublimate are dissolved in 50 or 60 of water (warmth quickens solution), and 5 of iodide of potassium in 15 or 20 of water, the solutions mixed, and the precipitate collected on a filter, drained, washed twice with distilled water, and dried on a plate over a water-bath. Mercuric nitrate, which is more soluble, and therefore somewhat more convenient for use on the large scale, may be used instead of the mercuric chloride. The mercury in mercuric or mercurous iodide is set free and sublimes in globules on heating either powder with dried carbonate of sodium in a test-tube; the iodine may be detected by digesting with solution of soda, filtering, and to the solution of iodide of sodium thus formed adding starch paste and acidulating with nitrous acid, when blue iodide of starch results. Mercuric iodide is insoluble in water, slightly soluble in alcohol, tolerably soluble in ether. Precipitated red iodide of mercury mixed with white wax, lard, and oil, forms the *Unguentum Hydrargyri Iodidi Rubri*, B. P. and U. S. P. *Donovan's Solution* contained mercuric and arsenious iodides.

## The Two Nitrates.

*Second Synthetical Reaction.*—Mix a little nitric acid in a test-tube with four or five times its bulk of water, add a small globule of mercury, and set the tube aside for a few hours, in a cool place; solution of mercurous nitrate ($HgNO_3$) will be

formed, and nitric oxide (NO) evolved. The solution may be retained for subsequent analytical operations.

$$Hg_3 + 4HNO_3 = 3HgNO_3 + 2H_2O + NO.$$

*Third Synthetical Reaction.*—Place mercury in strong nitric acid, and warm the mixture; mercuric nitrate is formed, and will be deposited in crystals as the solution cools. Retain the product for a subsequent experiment.

*The mercuric nitrates* vary somewhat in composition, according to the proportion, strength, and temperature of the acid used in their formation. A mercuric nitrate may be obtained having the formula $Hg2NO_3$.

| $Hg_3$ | + | $8HNO_3$ | = | $3(Hg2NO_3)$ | + | $2NO$ | + | $4H_2O$ |
|---|---|---|---|---|---|---|---|---|
| Mercury. | | Nitric acid. | | Mercuric nitrate. | | Nitric oxide. | | Water. |

*Mercuric Oxynitrates.*—From the normal mercuric nitrate several oxynitrates may be obtained. Thus on merely evaporating a solution of mercuric nitrate, and cooling, crystals having the formula $Hg_3O_36NO_3$ are deposited. The latter, by washing with cold water, yield a *yellow* pulverulent oxynitrate, $Hg_6O_4NO_3$: mixed with lard, this has sometimes been used as an ointment. Boiled in water, the yellow gives a *brick-red* oxynitrate, $Hg_6O_52NO_3$.

The pharmacopœial preparations of mercuric nitrate are *Liquor Hydrargyri Nitratis Acidus*, B. P. (sp. gr. 2.246; U. S. P., sp. gr. 2.165), and *Unguentum Hydrargyri Nitratis*, B. P. and U. S. P. The former (B. P.) is made by placing four ounces of mercury in five fluidounces of nitric acid diluted with an ounce and a half of water, and, when the metal is dissolved, boiling gently for fifteen minutes.

## The two Sulphates.

*Fourth Synthetical Reaction.*—Boil two or three grains of mercury with a few drops of strong sulphuric acid in a test-tube, or, better, small dish; sulphurous acid gas ($SO_2$) is evolved, and mercuric sulphate (*Hydrargyri Sulphas*, B. P.) ($HgSO_4$) results—a white heavy crystalline powder.

| $Hg$ | + | $2H_2SO_4$ | = | $HgSO_4$ | + | $SO_2$ | + | $2H_2O$ |
|---|---|---|---|---|---|---|---|---|
| Mercury. | | Sulphuric acid. | | Mercuric sulphate. | | Sulphurous acid gas. | | Water. |

Between two and three ounces of mercuric sulphate may be prepared from a fluidrachm of mercury and a fluidounce of sulphuric acid boiled together in a small dish. These are the official proportions. The operation is completed and any excess of acid removed by evaporating the mixture of metal and liquid to dryness, either in the open air or in a fume-chamber, sulphuric acid vapors being excessively irritating to the mucous membrane of the nose and throat; dry crystalline mercuric

sulphate remains. If residual particles of mercury are observed, the mass should be dampened with sulphuric acid and again heated.

*By-products.*—In chemical manufactories, secondary products, such as the sulphurous gas of the above reaction, are termed *by-products*, and, if of value, are utilized. In the present case the gas is of no immediate use, and is therefore allowed to escape. When very pure sulphurous acid gas is required for experiments on the small scale, this would be the best method of making it, a delivery-tube being adapted by a cork to the mouth of a flask containing the acid and metal. The sulphate of mercury would then become the by-product.

*Mercuric Oxysulphate.*—Water decomposes mercuric sulphate into a soluble acid salt and an insoluble yellow oxysulphate ($Hg_3O_2SO_4$). The latter is called *Turpeth mineral*, from its resemblance in color to the powdered root of *Ipomœa turpethum*, an Indian substitute for jalap. The yellow sulphate of mercury (*Hydrargyri Sulphas Flava*, U. S. P.) was formerly official in the Pharmacopœia of Great Britain, but is now seldom used.

*Fifth Synthetical Reaction.*—Rub a portion of the dry mercuric sulphate of the previous reaction with as much mercury as it already contains; the product, when the two have thoroughly blended, is mercurous sulphate ($Hg_2SO_4$): it may be retained for a subsequent experiment.

*Molecular Weight.*—The exact proportion of mercury to sulphate is merely a matter of calculation; for the combining proportion of a compound (if it possess any combining power) is the sum of the combining proportions of its constituents. *In other words, the combining weight of a molecule is simply the sum of the weights of its constituent atoms*, or more generally, *the molecular weight of a compound is the sum of the atomic weights of its elements.* In accordance with this rule (sometimes called the fourth law of chemical combination, though only a deduction from the first—p. 47), 296 of mercuric sulphate and 200 of mercury (about 3 to 2) are the exact proportions necessary to the formation of mercurous sulphate.

## The two Chlorides.

*Sixth Synthetical Reaction.*—Mix thoroughly a few grains of dry mercuric sulphate with about four-fifths its weight of chloride of sodium, and heat the mixture slowly in a test-tube in a fume-chamber or in the open air to leeward of the operator; mercuric chloride ($HgCl_2$), or *corrosive sublimate*, *Hydrargyri Perchloridum*, B. P., *Hydrargyri Chloridum Corrosivum*, U. S. P.), sublimes and condenses in the upper part of the tube in heavy colorless crystals or a crystalline mass.

Somewhat larger quantities (in the proportion of 20 of sulphate to 16 of salt, and, *vide infra*, 1 of black oxide of manganese) may be sublimed in a pair of two-ounce or three-ounce round-bottomed gallipots, the one inverted over the other, and the joint luted by moist fireclay (the powdered clay kneaded with water to the consistence of dough). The luting having been allowed to dry (somewhat slowly, to avoid cracks), the pots are placed upright on a sand-tray (plate-shape answers very well), sand piled round the lower and a portion of the upper pot, and the whole heated over a good-sized gas flame (if an air-gas flame, it should be an inch and a half wide and four or five inches long) for an hour or more (see fig. 36—pots raised to show joint). Red Iodide of Mercury and Calomel may be sublimed in the same way. The former requires less, the latter more, heat than corrosive sublimate.

Fig. 36.

Sublimation.

$$\underset{\text{Mercuric sulphate.}}{HgSO_4} + \underset{\text{Chloride of sodium.}}{2NaCl} = \underset{\text{Mercuric chloride.}}{HgCl_2} + \underset{\text{Sulphate of sodium.}}{Na_2SO_4}$$

*Note.*—If the mercuric sulphate contain any mercurous sulphate, some calomel may be formed. This result will be avoided if 2 or 3 per cent. of black oxide of manganese be previously mixed with the ingredients, the action of which is to eliminate chlorine from the excess of chloride of sodium used in the process, the chlorine converting any calomel into corrosive sublimate. Manganate of sodium and a lower oxide of manganese are simultaneously produced.

*Precaution.*—The operation is directed to be conducted with care in a fume-chamber or in the open air, because the vapor of corrosive sublimate, which might possibly escape, is very acrid and highly poisonous. Its vulgar name is indicative of its properties.

Ten grains of perchloride of mercury and the same quantity of chloride of ammonium in one pint of water form the *Liquor Hydrargyri Perchloridi*, B. P. A dilute aqueous solution of perchloride of mercury is liable to decomposition, calomel being precipitated, water decomposed, hydrochloric acid formed, and oxygen gas evolved. The presence of excess of chloride of ammonium, with a portion of which the mercuric chloride forms a stable double salt, prevents the decomposition.

*Seventh Synthetical Reaction.*—Mix a few grains of the mercurous sulphate of the fifth reaction with about a third of

its weight of chloride of sodium, and sublime in a test-tube; crystalline mercurous chloride (HgCl) or calomel (*Hydrargyri Subchloridum*, B. P., *Hydrargyri Chloridum Mite*, U. S. P.) results. Larger quantities may be prepared in the manner directed for corrosive sublimate, a somewhat higher temperature being employed; similar precaution must also be observed. The proportions are 10 of mercuric sulphate to 7 of mercury and 5 of dry chloride of sodium. "Moisten the sulphate of mercury with some of the water, and rub it and the mercury together until globules are no longer visible; add the chloride of sodium, and thoroughly mix the whole by continued trituration. When dry sublime by a suitable apparatus into a chamber of such a size that the calomel, instead af adhering to its sides as a crystalline crust, shall fall as a fine (dull-white) powder on its floor. Wash this powder with boiling distilled water until the washings cease to be darkened by a drop of sulphydrate of ammonium. Finally, dry at a heat not exceeding 212° F., and preserve in a jar or bottle impervious to light."

$$\underset{\text{Mercurous sulphate.}}{Hg_2SO_4} + \underset{\text{Chloride of sodium.}}{2NaCl} = \underset{\text{Mercurous chloride.}}{2HgCl} + \underset{\text{Sulphate of sodium.}}{Na_2SO_4}$$

*The term calomel* (καλός, *kalos*, good, and μέλας, *melas*, black) is said to relate to the use of the salt as a *good* remedy for *black* bile, but probably was simply indicative of the esteem in which black sulphide of mercury was held, the compound to which the name calomel was first applied.

*Test for corrosive sublimate in calomel.*—If the mercurous sulphate contains mercuric sulphate, some mercuric chloride will also be formed. Corrosive sublimate is soluble in water, calomel insoluble; the presence of the former may therefore be proved by boiling a few grains of the calomel in distilled water, filtering and testing by sulphuretted hydrogen or sulphydrate of ammonium as described hereafter. If corrosive sublimate is present the whole bulk of the calomel must be washed with hot distilled water till the filtrate ceases to give any indications of mercury. Corrosive sublimate is more soluble in alcohol, and still more in ether; calomel insoluble. Ether in which calomel has been digested should, therefore, after filtration, yield no residue on evaporation. Calomel is converted by hydrocyanic acid into mercuric salt, and a black powder readily yielding metallic mercury. Powell and Bayne have shown that a certain proportion of hydrochloric acid arrests this action.

*Note.*—The above process is that of the Pharmacopœia; but calomel may also be made by other methods. Calomel mixed with lard forms the *Unguentum Hydrargyri Subchloridi*, B. P., and with sulphurated antimony, guaiacum, resin, and castor oil, the *Pilula Hydrargyri Subchloridi Composita*, B. P., *Pilula Antimonii Composita*, U. S. P., or "Plummer's Pills."

## The Two Oxides.

*Eighth Synthetical Reaction.*—Evaporate the mercuric nitrate of the third reaction to dryness in a small dish, in a fume-chamber, or in the open air if more than a few grains have been prepared, and heat the residue till no more fumes are evolved; mercuric oxide (HgO), "Red Precipitate," the Red Oxide of Mercury (*Hydrargyri Oxidum Rubrum*, B. P. and U. S. P.), remains.

$$\underset{\text{Mercuric nitrate.}}{2(Hg2NO_3)} = \underset{\text{Mercuric oxide.}}{2HgO} + \underset{\text{Nitric peroxide.}}{4NO_2} + \underset{\text{Oxygen.}}{O_2}$$

The nitric constituents of the salt may be partially economized by previously thoroughly mixing with the dry mercuric nitrate as much mercury as is used in its preparation, or as much as it already contains (ascertained by calculation from the atomic weights and the weight of nitrate under operation, as in making mercurous sulphate p. 200), and well heating the mixture. In this case the free mercury is also converted into mercuric oxide. This is the official process, the Pharmacopœial quantities being four ounces of mercury dissolved in four and a half fluidounces of nitric acid diluted with two ounces of water, the solution evaporated to dryness, the residue thoroughly mixed with four ounces of mercury, and the whole heated until acid vapors cease to be evolved. (Mercuric oxide is tested for nitrate by heating a little of the sample in a test-tube, when orange nitrous vapors are produced and are visible in the upper part of the tube, if nitrate is present.)

$$\underset{\text{Mercuric nitrate.}}{Hg2NO_3} + \underset{\text{Mercury.}}{Hg} = \underset{\text{Mercuric oxide.}}{2HgO} + \underset{\text{Nitric peroxide.}}{2NO_2}$$

*Mercuric oxide* is an orange-red powder, more or less crystalline according to the extent to which it may have been stirred during preparation from the nitrate, much rubbing giving the crystals a pulverulent character. Mixed with yellow wax and oil of almonds it yields the *Unguentum Hydrargyri Oxidi Rubri*, B. P. (1 part in 8). A similar ointment is official in U. S. P. Mercuric oxide, in contact with oxidizable organic matter, is liable to reduction to black or mercurous oxide.

*Ninth Synthetical Reaction.*—To solution of potash or soda, or lime-water, in a test-tube or larger vessel, add solution of corrosive sublimate or of mercuric nitrate; yellow oxide of mercury, *Hydrargyri Oxidum Flavum*, B. P. and U. S. P., or mercuric oxide (HgO), is precipitated.

$$\underset{\text{Mercuric chloride.}}{HgCl_2} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Mercuric oxide.}}{HgO} + \underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Water.}}{H_2O}$$

Eighteen grains of corrosive sublimate to ten ounces of lime-water form the *Lotio Hydrargyri Flava*, B. P. The precipitate only differs physically from the red mercuric oxide; the yellow is in a more minute state of division than the red. Mercuric oxide is very slightly soluble in water, but sufficiently so to communicate a decidedly metallic taste.

*Tenth Synthetical Reaction.*—To calomel add solution of potash or soda, or lime-water; black oxide of mercury, or mercurous oxide ($Hg_2O$) is produced, and may be filtered off, washed, and dried. (This reaction and the formation of a white curdy precipitate, on the addition of solution of nitrate of silver to the filtrate from the mercurous oxide, acidified by nitric acid, form sufficient evidence of a powder being or containing calomel. The curdy precipitate is chloride of silver.)

Thirty grains of calomel to ten ounces of lime-water form the *Lotio Hydrargyri Nigra*, B. P.

$$\underset{\text{Mercurous chloride.}}{2HgCl} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Mercurous oxide.}}{Hg_2O} + \underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Water.}}{H_2O}$$

### (*b*) *Analytical Reactions* (*Tests*).

(The mercury occurring as mercuric or mercurous salts.)

*First Analytical Reaction.—The Copper Test.*—Deposition of mercury upon, and sublimation from copper.—Place a small piece of bright copper, about half an inch long and a quarter of an inch broad, in a solution of any salt of mercury, mercurous or mercuric, and heat in a test-tube; the copper becomes coated with mercury in a fine state of division. (The absence of any notable quantity of nitric acid must be insured or the copper itself will be dissolved. See next page.) Pour away the supernatent liquid from the copper, wash the latter once or twice by pouring water into, and then out of, the tube, remove the metal, take off excess of water by gentle pressure in a piece of filter-paper, dry the copper by passing it quickly through a flame, holding it by the fingers; finally, place the copper in a dry narrow test-tube, and heat to redness in a flame, the tube being held in a horizontal position; the mercury sublimes and condenses as a white sublimate of minute globules on the cool part of the tube outside the flame. The globules aggregate on gently pressing with a glass rod, and are especially visible where flattened between the rod and the side of the test-tube.

*Notes on the Test.*—This is a valuable test, for several reasons: It is very delicate when performed with care. It brings before the

observer the element itself—one which from its metallic lustre and fluidity cannot be mistaken for any other. It separates the element both from mercurous and mercuric salts. Mercury can in this way readily be eliminated in the presence of most other substances, organic or inorganic.

In performing the test the presence of any quantity of nitric acid may be avoided by adding an alkali until a slight permanent precipitate appears, and then reacidifying with a few drops of acetic or hydrochloric acid; or by concentrating in an evaporating-dish after adding a little sulphuric acid, and then rediluting.

*Tests continued.* (The mercury occurring as mercuric salt.)

*Second Analytical Reaction.*—To a few drops of a solution of a mercuric salt (corrosive sublimate, for example) add solution of iodide of potassium, drop by drop; a precipitate of mercury iodide ($HgI_2$) forms, and at first quickly redissolves, but is permanent when sufficient iodide of potassium has been added. Continue the addition of iodide of potassium; the precipitate is once more redissolved.

*Note.*—When first precipitated, mercuric iodide is yellowish-red, but soon changes to a beautiful scarlet. Its solubility either in solution of the mercuric salt or in solution of iodide of potassium renders the detection of a small quantity of a mercuric salt by iodide of potassium, or a small quantity of an iodide by a mercuric solution, difficult, and hence lessens the value of the reaction as a test.

*Third Analytical Reaction.*—Add a solution of mercuric salt to solution of ammonia, taking care that the mixture, after well stirring, still smells of ammonia; a white precipitate falls.

## Ammoniated Mercury.

Performed in a test-tube, this reaction is a very delicate test of the presence of a mercuric salt; performed in larger vessels, the mercuric salt being corrosive sublimate (3 ounces dissolved in 3 pints of distilled water, the solution poured into 4 fluidounces of Solution of Ammonia, and the precipitate washed and dried over a water-bath), it is the usual process for the preparation of "white precipitate," the old "ammonia chloride," or "amido-chloride of mercury" (so called because then considered to be a compound of mercury with chlorine and with *amidogen*, $NH_2$, or $HgCl_2$, $Hg2NH_2$, chloride and *amide* of mercury), now known as Ammoniated Mercury (*Hydrargyrum Ammoniatum*, B. P. and U. S. P.).

*Constitution of Ammoniated Mercury.*—This precipitate is considered to be the chloride of mercuric ammonium —$NH_2Hg''Cl$— that is, chloride of ammonium ($NH_4Cl$) in which two univalent atoms of hydrogen are replaced by one bivalent atom of mercury.

$$\underset{\text{Mercuric chloride.}}{HgCl_2} + \underset{\text{Ammonia.}}{2NH_4HO} = \underset{\text{"White precipitate."}}{NH_2Hg''Cl} + \underset{\text{Chloride of ammonium.}}{NH_4Cl} + \underset{\text{Water.}}{2H_2O}$$

*Varieties of Ammoniated Mercury.*—If the order of mixing be reversed and ammonia be added to solution of mercuric chloride, a double chloride of mercuric-ammonium and mercury results ($NH_2HgCl,HgCl_2$): it contains 76.55 per cent. of mercury. Previously to the year 1826, "white precipitate" was officially made by adding a fixed alkali to a solution of equal parts of corrosive sublimate and sal-ammoniac; this gave a double chloride of mercuric ammonium and ammonium ($NH_2HgCl,NH_4Cl$), containing 65.57 per cent. of mercury. This compound is now known as "*fusible* white precipitate," because at a temperature somewhat below redness it fuses and then volatilizes. The "white precipitate" which has been official since 1826 contains 79.52 per cent. of mercury. The true compound may be distinguished as "*infusible* white precipitate," from the fact that when heated it volatilizes without fusing. An ointment of this body is official (*Unguentum Hydrargyri Ammoniati*, B. P. and U. S. P.). Prolonged washing with water converts "white precipitate" into a yellowish compound ($NH_2HgCl,HgO$); hence the official preparation is seldom thoroughly freed from the chloride of ammonium which is formed during its manufacture; and which, if present in larger proportion than seven or eight per cent., gives to it the character of partial or complete fusibility. With iodine, chlorine, or bromine, white precipitate may yield the highly explosive iodide, chloride, or bromide of nitrogen.

*Note.*—Chloride of mercuric-ammonium is only one member of a large class of similar compounds, derivable from the various salts of ammonium by displacement of atoms of hydrogen by other atoms. The composition of the chloride of mercurous-ammonium (see next page) and of ammonio-nitrate of silver is consistent with this view.

$$N\left\{\begin{matrix}H\\H\\H\\H\end{matrix}\right\}Cl \qquad N\left\{\begin{matrix}Hg'\\Hg'\\H\\H\end{matrix}\right\}Cl \qquad N\left\{\begin{matrix}Hg''\\H\\H\end{matrix}\right\}Cl \qquad N\left\{\begin{matrix}Ag\\Am\\H\\H\end{matrix}\right\}NO_3$$

Chloride of common ammonium. | Chloride of mercurous-ammonium. | Chloride of mercuric-ammonium. | Nitrate of argent-ammon-ammonium.

The composition of the ammonio-sulphates of copper (pp. 177 and 193) is consistent with the second and third of the following formulæ, the first being that of sulphate of ammonia:—

$$N_2\left\{\begin{matrix}H_2\\H_2\\H_2\\H_2\end{matrix}\right\}SO_4 \qquad N_2\left\{\begin{matrix}Cu''\\Am_2\\H_2\\H_2\end{matrix}\right\}SO_4 \qquad N_2\left\{\begin{matrix}Cu''\\Am_2\\Am_2\\H_2\end{matrix}\right\}SO_4$$

The iodide of dimercuric ammonium ($NHg''_2I$) is formed in testing for ammonia by the "Nessler" reagent (*vide* Index).

*Fourth Analytical Reaction.*—Pass sulphuretted hydrogen through a mercuric solution; a black precipitate of mercuric sulphide (HgS) falls.

*Note.*—Sulphuretted hydrogen also precipitates mercurous sulphide ($Hg_2S$) from mercurous solutions; and in appearance the precipitates are alike; hence this reagent does not distinguish between mercurous and mercuric salts. But in the course of systematic analysis, mercuric salts are thrown down from solution as sulphide after mercurous salts have been otherwise removed. The sulphides are insoluble in sulphydrate of ammonium.

*Note.*—An insufficient amount of the gas gives a white or colored precipitate of oxysulphide. Prolonged contact with sulphuretted hydrogen-water or a sulphydrate, especially when the mixture is kept warm, converts the black into a red sulphide.

*Æthiops Mineral*, the *Hydrargyri Sulphuretum cum Sulphure*, is a mixture of sulphide of mercury and sulphur, obtained on triturating the elements in a mortar till globules are no longer visible. Its name is probably in allusion to its similarity in color to the skin of the Æthiop. It was formerly official.

*Vermilion* is mercuric sulphide prepared by sublimation (*Hydrargyri Sulphuretum Rubrum*, U. S. P.).

*Tests continued.* (The mercury occurring as mercurous salt.)

*Fifth Analytical Reaction.*—To a solution of a mercurous salt (the mercurous nitrate obtained in the second synthetical reaction, for example) add hydrochloric acid, or any soluble chloride; a white precipitate of calomel (HgCl) occurs.

This reaction was formerly official in the Dublin Pharmacopœia as a process for the preparation of calomel.

*Sixth Analytical Reaction.*—To solution of a mercurous salt add iodide of potassium; green mercurous iodide (HgI) is precipitated.

*Seventh Analytical Reaction.*—To a mercurous salt, dissolved or undissolved (calomel), add ammonia; black salt (chloride) of mercurous ammonium ($NH_2Hg_2Cl$) is formed (see previous page).

### *Other Tests for Mercury.*

The elimination of mercury in the actual state of metal by the copper test, coupled with the production or non-production of a white precipitate on the addition of hydrochloric acid to the original solution, is usually sufficient evidence of the presence of mercury and its existence as a mercurous or mercuric salt. But other tests may sometimes be applied with advantage. Thus, metallic mercury is deposited on placing a drop of the solution on a plate of gold (sovereign or half sovereign), and touching the drop and the edge of the plate simultaneously with a key; an electric current passes, under these circumstances, from the gold to the key, and thence through the

liquid to the gold, decomposing the salt, the mercury of which forms a white metallic spot on the gold, while the other elements go to the iron. This is called *the galvanic test*, and is useful for clinical purposes.——Solution of stannous chloride ($SnCl_2$)—see Index—from the readiness with which it forms stannic salts ($SnCl_4$, $SnO_2$, etc.), gives a white precipitate of mercurous chloride in mercuric solutions, and quickly still further reduces this mercurous chloride (and other mercury-salts) to a grayish mass of finely divided mercury; this is the old *magpie test*, probably so called from the white and gray appearance of the precipitate. The reaction may even be obtained from such insoluble mercury compounds as "white precipitate."——Confirmatory tests for mercuric and mercurous salts will be found in the action of solution of potash, solution of soda, lime-water, solution of ammonia, and solution of iodide of potassium. (*Vide* pages 203 to 207.)——Normal alkaline carbonates produce yellowish mercurous carbonate, and brownish-red mercuric carbonate, both of them unstable.——Alkaline bicarbonates give mercurous carbonate, with mercurous salts, and with mercuric salts white (becoming red) mercuric oxysalt. —— Yellow chromate of potassium ($K_2CrO_4$) gives, with mercurous salts, a red precipitate of mercurous chromate ($Hg_2CrO_4$).——Mercury and all its compounds are volatile, many of them being decomposed, at the same time yielding globules of condensed metal: the experiment is most conveniently performed in a test-tube.

*Antidote.*—Albumen gives a white precipitate with solution of mercuric salts; hence the importance of administering white of egg while waiting for a stomach-pump in cases of poisoning by corrosive sublimate.

---

## QUESTIONS AND EXERCISES.

293. Name the chief ore of mercury, and describe a process for the extraction of the metal.

294. Give the properties of mercury.

295. In what state does mercury exist in "Gray Powder?"

296. What other preparations of metallic mercury itself are employed in medicine?

297. State the relation of the mercurous to the mercuric compounds.

298. Distinguish between an *alloy* and an *amalgam*.

299. State the formulæ of the two Iodides of Mercury.

300. Under what circumstances does mercuric iodide assume two different colors?

301. Illustrate the chemical law of Multiple Proportions as explained by the atomic theory, employing for that purpose the stated composition of the two iodides of mercury.

302. Write down the formulæ of Mercurous and Mercuric Nitrates and Sulphates

303. How is Mercuric Sulphate prepared?

304. What is the formula of "Turpeth Mineral?"

305. Describe the processes necessary for the conversion of mercury into Calomel and Corrosive Sublimate, using diagrams.

306. Why is black oxide of manganese sometimes mixed with the other ingredients in the preparation of Corrosive Sublimate?

307. Give the chemical and physical points of difference between Calomel and Corrosive Sublimate.

308. How may a small quantity of Calomel in Corrosive Sublimate be detected?

309. Work out a sum showing how much mercury will be required in the manufacture of one ton of Calomel. *Ans.* 17 cwt. nearly.

310. Mention official preparations of the chlorides of mercury.

311. Give the formulæ and mode of formation of the Red, Yellow, and Black Oxides of Mercury, employing diagrams.

312. Explain the action of the chief general test for mercury.

313. How are mercurous and mercuric salts analytically distinguished?

314. Give a probable view of the constitution of *Hydrargyrum Ammoniatum*, and an equation showing how it is made.

315. What is the best temporary antidote in cases of poisoning by mercury?

---

## LEAD.

Symbol Pb. Atomic weight 207.

*Source.*—The ores of lead are numerous; but the one from which the metal is chiefly obtained is the sulphide of lead (PbS), or *galena* (from γαλήνη, *galene*, tranquillity, perhaps from its supposed effect in allaying pain).

*Preparation.*—The ore is first roasted in a current of air; much sulphur is thus burnt off as sulphurous acid gas, while some of the metal is converted into oxide and a portion of the sulphide oxidized to sulphate. Oxidation being stopped when the mass presents certain appearances, the temperature is raised, and the oxide and sulphate, reacting on undecomposed sulphide, yield the metal and much sulphurous acid gas:—

$$2PbO + PbS = Pb_3 + SO_2$$
$$PbSO_4 + PbS = Pb_2 + 2SO_2.$$

*Uses.*—The uses of lead are well known. Alloyed with arsenicum it forms common *shot*, with antimony gives *type-metal*, with tin *solder*, and in smaller quantities enters into the composition of *Britannia metal*, *pewter*, and other alloys. Lead is so slightly attacked by

acids that chemical vessels and instruments are often made of it. Even hot hydrochloric acid only slowly converts it into chloride of lead, with evolution of hydrogen. Sulphuric acid by aid of air only very slowly attacks it, with formation of sulphate of lead and water. Even nitric acid very slowly converts it into nitrate, with evolution of nitric oxide and nitrous oxide gases and water.

*The salts of lead* used in pharmacy and all other preparations of lead are obtained, directly or indirectly, from the metal itself. Heated in a current of air, lead combines with oxygen and forms oxide of lead (PbO) (*Plumbi Oxidum*, B. P. and U. S. P.), a yellow powder (*massicot*), or, if fused and solidified, a brighter reddish-yellow heavy mass of bright scales, termed *litharge* (from λίθος, *lithos*, a stone, and ἀργυρος, *argyros*, silver). It is from this oxide that the chief lead compounds are obtained. Oxide of lead, by further roasting in a current of air, yields *red lead* (or *minium*), $Pb_3O_4$, or $PbO_2 2PbO$. Both oxides are much used by painters, paper-stainers, and glass-manufacturers. *White lead* is a mixture of carbonate ($PbCO_3$) and hydrate of lead (Pb2HO) (commonly 2 molecules of the former to 1 of the latter), usually ground up with about 7 per cent. of linseed oil; it is made by exposing lead, cast in spirals or little gratings, to the action of air, acetic fumes, and carbonic acid, the latter generated from decaying vegetable matter, such as spent tan; oxyacetate of lead slowly but continuously forms, and is as continuously decomposed by the carbonic acid with production of hydrate and carbonate, or *dry white lead*. The grating-like masses, when ground, form the heavy white pulverulent official *Plumbi Carbonas*, B. P. and U. S. P. The latter is the active constituent of *Unguentum Plumbi Carbonatis*, B. P. and U. S. P., the old *Unguentum Cerussæ*.

*Lead compounds are poisonous*, producing saturnine colic, or even paralysis. These effects are termed *saturnine* from an old name of lead, *Saturn*. The alchemists called lead Saturn, first, because they thought it the oldest of the seven then known metals, and it might therefore be compared to Saturn, who was supposed to be the father of the gods; and, secondly, because its power of dissolving other metals recalled a peculiarity of Saturn, who was said to be in the habit of devouring his own children.

*Quantivalence.* — The atom of lead is sometimes quadrivalent ($Pb''''$); but in most of the compounds used in medicine it exerts bivalent activity only ($Pb''$).

## Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

### (*a*) *Synthetical Reactions.*

### Acetate of Lead.

*First Synthetical Reaction.*—Place a few grains of oxide of lead in a test-tube, add about an equal weight of water and two and a half times its weight of acetic acid, and boil; the oxide dissolves (or, rather, disappears—dissolves with simul-

taneous decomposition) and forms a solution of acetate of lead ($Pb2C_2H_3O_2$). When cold, or on evaporation (the solution being kept faintly acid), crystals of acetate of lead ($Pb2C_2H_3O_2$, $3H_2O$) are deposited. Larger quantities are obtained by the same method.

$$\underset{\text{Oxide of lead.}}{PbO} + \underset{\text{Acetic acid.}}{2HC_2H_3O_2} = \underset{\text{Acetate of lead.}}{Pb2C_2H_3O_2} + \underset{\text{Water.}}{H_2O}$$

This is the official process for *Plumbi Acetas*, B. P. and U. S. P. The salt is termed *Sugar of Lead*, from its sweet taste. Besides its direct use in Pharmacy, it forms three-fourths of the *Pilula Plumbi cum Opio*, B. P., is the chief constituent of *Unguentum Plumbi Acetatis*, B. P., and an ingredient in *Suppositoria Plumbi Composita*, B. P., and *Suppositoria Plumbi* and *Suppositoria Plumbi et Opii*, U. S. P.

## Subacetate or Oxyacetate of Lead.

*Second Synthetical Reaction.*—Boil acetate of lead with about four times its weight of water, and rather more than two-thirds its weight of oxide of lead; the resulting filtered liquid is solution of oxyacetate of lead, *Liquor Plumbi Subacetatis*, B. P. and U. S. P.

The official (B. P.) *Liquor* is made by boiling 5 ounces of acetate and 3½ of oxide in one pint of distilled water for half an hour (constantly stirring), filtering, and making up for any loss by evaporation by diluting the filtrate to 1 pint.

A similar solution was used by M. Goulard, who called it *Extractum Saturni*, and drew attention to it in 1770. It is now frequently termed *Goulard's Extract*. A more dilute solution, 1 of *Liquor* and 1 of spirit in 80 of *distilled* water, is also official in the Pharmacopœias, under the name of *Liquor Plumbi Subacetatis Dilutus*. The latter is commonly known as *Goulard Water*. The stronger solution is the chief ingredient in *Unguentum Plumbi Subacetatis Compositum*, B. P., a slight modification of the old *Goulard's Cerate*. Similar preparations are official in the United States Pharmacopœia.

*Oxyacetates of Lead.*—The official subacetate of lead is not a definite chemical salt. It is probably a mixture of two subacetates of lead, which are well-known crystalline compounds, and which the author is disposed to regard as having a constitution similar to that he has already indicated for some other salts (see Iron, Antimony, and Bismuth). Exposed to air it absorbs carbonic acid gas, and hydrato-carbonate of lead is deposited.

B. P.
- Acetate of Lead (3 molecules) . . $Pb_3\ 6C_2H_3O_2$
- Pyro-oxyacetate of lead . . . . . $Pb_3O4C_2H_3O_2$
- Goulard's oxyacetate of lead . . . $Pb_3O_2 2C_2H_3O_2$
- Oxide of lead (3 molecules) . . . . $Pb_3O_3$.

$$\underset{\text{Oxide of lead.}}{PbO} + \underset{\text{Acetate of lead.}}{Pb2C_2H_3O_2} = \underset{\text{Official "subacetate."}}{Pb_3O2C_2H_3O_2}$$

$$\text{or } \underset{\text{Oxide of lead.}}{3PbO} + \underset{\text{Acetate of lead.}}{3(Pb2C_2H_3O_2)} = \underset{\text{Pyro-oxyacetate.}}{Pb_3O4C_2H_3O_2} + \underset{\text{Goulard's oxyacetate. The official "subacetate."}}{Pb_3O_2C_2H_3O_2}$$

## Nitrate of Lead. Red Lead. Peroxide of Lead.

*Third Synthetical Reaction.* Digest a few grains of red lead in nitric acid and water; nitrate of lead ($Pb2NO_3$) is formed, and remains in solution, while a puce-colored peroxide of lead ($PbO_2$) is precipitated.

*Nitrate of Lead* (*Plumbi Nitras*, B. P. and U. S. P.) is more directly made by dissolving litharge (PbO) in nitric acid—

$$PbO + 2HNO_3 = Pb2NO_3 + H_2O;$$

but the former reaction serves to bring before the reader two other oxides of lead, namely, red lead ($Pb_3O_4$) and peroxide of lead ($PbO_2$). In the latter oxide the quadrivalent character of lead is obvious. Nitrate of lead is used officially in preparing iodide of lead; for this purpose the above mixture is filtered, the precipitate of peroxide of lead purified from adhering nitrate by passing hot water through the filter, the filtrate and washings evaporated to dryness to remove excess of nitric acid, the residual nitrate of lead redissolved by ebullition with a small quantity of hot water, and the solution set aside to crystallize, or a portion at once used for the following experiment. Nitrate of lead forms white crystals derived from octahedra.

## Iodide of Lead.

*Fourth Synthetical Reaction.*—To a neutral solution of nitrate of lead add solution of iodide of potassium; a precipitate of iodide of lead ($PbI_2$) falls (*Plumbi Iodidum*, B. P. and U. S. P.). Equal weights of the salts may be used in making large quantities.

$$\underset{\text{Nitrate of lead.}}{Pb2NO_3} + \underset{\text{Iodide of potassium.}}{2KI} = \underset{\text{Iodide of lead.}}{PbI_2} + \underset{\text{Nitrate of potassium.}}{2KNO_3}$$

Iodide of lead is the chief ingredient in *Emplastrum Plumbi Iodidi*, B. P., and *Unguentum Plumbi Iodidi*, B. P. and U. S. P. The iodide of lead used in making the plaster is altered in the process, either iodide of sodium being formed (from the hard soap present), or possibly a compound of the iodide and oleate of lead, corresponding with the almost colorless hydrato-iodides of lead.

*Crystals of Iodide of Lead.*—Heat the iodide of lead with the supernatant liquid, and if necessary filter; the salt is dissolved, and again separates in golden crystalline scales as the solution cools.

## Oleate of Lead (Lead Plaster).

*Fifth Synthetical Reaction.*—Boil together in a small dish some very finely-powdered oxide of lead, with about twice its weight of olive oil, and ten or twenty times as much water, well stirring the mixture, and from time to time replacing water that has evaporated; the product is a white mass of oleate of lead ($Pb2C_{18}H_{33}O_2$) (*Emplastrum Plumbi*, B. P. and U. S. P.), glycerine remaining in solution in the water. Larger quantities are prepared in the same manner.

$$\underset{\text{Oxide of lead.}}{3PbO} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Oleate of glyceryl (olive-oil or oleine).}}{2(C_3H_53C_{18}H_{33}O_2)} =$$

$$\underset{\text{Oleate of lead (lead plaster).}}{3(Pb2C_{18}H_{33}O_2)} + \underset{\text{Hydrate of glyceryl (glycerine).}}{2(C_3H_53HO)}$$

The action between the oxide of lead and olive oil is slow, requiring several hours for its completion.

*The glycerine* may be obtained by treating the aqueous product of the above reaction with sulphuretted hydrogen to remove a trace of lead, then digesting with animal charcoal, filtering and evaporating. But on the large scale glycerine is now usually produced as a by-product in the manufacture of candles; for its elements are found in all vegetable and animal fats. (*Vide* Index.) If in making lead plaster the mixture be evaporated to dryness, some of the glycerine will escape with the steam and some remain with the plaster.

---

Modes of forming *chloride*, *sulphide*, *chromate*, *sulphate*, *hydrate*, and other salts of lead are incidentally described in the following analytical paragraphs.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—To a solution of lead salt (acetate, for example) add hydrochloric acid; a white precipitate of chloride of lead ($PbCl_2$) is obtained. Boil the precipitate with much water; it dissolves, but, on the solution cooling, is redeposited in small acicular crystals. Filter the cold solution, and pass sulphuretted hydrogen through it; a black precipitate (sulphide of lead, PbS) shows that the chloride of lead is soluble to a slight extent in cold water.

*Note.*—A white precipitate on the addition of hydrochloric acid, soluble in hot water, and blackened by sulphuretted hydrogen, sufficiently distinguishes lead salts from those of other metals, but the non-production of such a precipitate does not prove the absence of

a small quantity of lead, chloride of lead being slightly soluble in cold water. Hydrochloric acid will be found to be a useful but not a delicate test for lead.

*Second Analytical Reaction.*—Through a dilute solution of a lead salt acidulated with hydrochloric acid pass sulphuretted hydrogen; a black precipitate of sulphide of lead (PbS) occurs.

*Lead in Water.*—The foregoing is a very delicate test. Should a trace of lead be present in water used for drinking purposes, sulphuretted hydrogen will detect it. On passing the gas through a pint of such acidulated water, a brownish color is produced. If the tint is scarcely perceptible, set the liquid aside for a day; the gas will become decomposed and a thin layer of sulphur be found at the bottom of the vessel, white if no lead be present, but more or less brown if it contain sulphide of lead.

*Third Analytical Reaction.*—To solution of a lead salt add sulphydrate of ammonium; a black precipitate of sulphide of lead falls, insoluble in excess.

*Fourth Analytical Reaction.*—To solution of a lead salt add solution of chromate of potassium ($K_2CrO_4$); a yellow precipitate of chromate of lead ($PbCrO_4$) is formed, insoluble in weak acids.

*Chromes.*—This reaction has technical as well as analytical interest. The precipitate is the common pigment termed *chrome yellow*, or *lemon chrome*. Boiled with lime and water, a portion of the chromic radical is removed as soluble chromate of calcium, and an oxychromate of lead, of a bright red or orange color (*orange chrome*), is produced.

*Fifth Analytical Reaction.*—To solution of a lead salt add dilute sulphuric acid, or a solution of a sulphate; a white precipitate of sulphate of lead ($PbSO_4$) falls.

*Sulphate of lead* is slightly soluble in strong acids, and in solutions of alkaline salts; it is insoluble in acetic acid. It is readily dissolved and indeed decomposed by solution of acetate of ammonium, the liquid yielding the ordinary reactions with soluble chromates and iodides.

In dilute solutions the above sulphuric reaction does not take place immediately; the precipitate, however, falls after a time; its appearance may be hastened by evaporating the solution nearly to dryness and then rediluting.

The white precipitate always noticed in the vessels in which diluted sulphuric acid is kept, is sulphate of lead, derived from the leaden chambers in which the acid is made; solubility in strong acid and insolubility in weak, explains its appearance.

*Antidotes.*—From the insolubility of sulphate of lead in water, the best *antidote*, in a case of poisoning by the acetate or other soluble salt of lead, is a soluble sulphate, such as Epsom salt, sulphate of

sodium or alum, vomiting being also induced, or the stomach-pump applied as quickly as possible.

*Other tests for lead* will be found in the reaction with *iodide of potassium* (*vide* p. 212); with *alkaline carbonates*, a white precipitate ($2PbCO_3 + Pb2HO$) insoluble in excess; with *alkalies*, a white precipitate ($Pb2HO$) more or less soluble in excess; with alkaline *phosphates*, *arseniates*, *ferrocyanides* and *cyanides*, precipitates mostly insoluble but of no special analytical interest. Insoluble salts of lead are decomposed by solutions of potash (KHO) or soda (NaHO).

The metal is precipitated in a beautifully crystalline state by metallic zinc and some other metals; the *lead tree* is thus formed.——The *blowpipe flame* decomposes solid lead compounds placed in a small cavity in a piece of charcoal, a soft malleable bead of metal being produced, and a yellowish ring of oxide deposited on the charcoal.

---

## QUESTIONS AND EXERCISES.

316. Write down equations descriptive of the smelting of galena.
317. Mention some of the alloys of lead.
318. How is litharge produced?
319. Give the formulæ of white lead and red lead.
320. Describe the manufacture of white lead.
321. What is the quantivalence of lead?
322. Draw a diagram expressive of the formation of Acetate of Lead.
323. Describe the preparation and composition of *Liquor Plumbi Subacetatis*.
324. What is the action of nitric acid on red lead, litharge, and metallic lead?
325. How is the official Iodide of Lead prepared?
326. Describe the reaction between oxide of lead, water, and olive oil, at the temperature of boiling water, and give chemical formulæ explanatory of the constitution of the products.
327. Mention the chief tests for lead.
328. How would you search for lead in potable water?
329. What is the composition of chrome yellow?
330. State a method whereby lead, barium, and silver may be separated.
331. Name the best antidote in case of poisoning by salts of lead.

## SILVER.

Symbol Ag. Atomic weight 108.

*Source.*—This element occurs in nature in the free state and as ore, the common variety being sulphide of silver ($Ag_2S$) in combination with much sulphide of lead, forming *argentiferous galena.*

*Preparation.*—The lead from such galena (p. 209) is melted and slowly cooled; crystals of lead separate and are raked out from the still fluid mass, and thus an alloy very rich in silver is finally obtained; this is roasted in a current of air, whereby the lead is oxidized and removed as litharge, pure silver remaining. Other ores undergo various preparatory treatments according to their nature, and are then shaken with mercury, which amalgamates with and dissolves the particles of silver, the mercury being subsequently removed from the amalgam by distillation. Soils and minerals containing metallic silver are also treated in this way. An important improvement in the amalgamation process, by which the mercury more readily unites with the silver, consists in the addition of a small proportion of sodium to the mercury—a discovery simultaneously made in England by Crookes, and in New York by Wurtz. Silver is not readily affected by the weak acids or other fluids of food, though it is rapidly tarnished by sulphur or sulphur compounds. It does not perceptibly attack hydrochloric acid; reduces strong nitric acid to nitrous anhydride ($N_2O_3$), and a weaker acid to nitric oxide (NO); it reduces hot sulphuric acid to sulphurous anhydride ($SO_2$), sulphate of silver ($Ag_2SO_4$) being formed. The latter salt is crystalline and slightly soluble in water.

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Synthetical Reactions.*

#### Impure Nitrate of Silver.

*First Synthetical Reaction.*—Dissolve a silver coin in nitric acid; nitric oxide gas (NO) and nitrous anhydride ($N_2O_3$) are evolved, and a solution of nitrates of silver and copper obtained.

*Silver Coinage.*—Pure silver is too soft for use as coin, it is therefore hardened by alloying with copper. The silver money of England contains 7.5, of Prussia 25 and of France 10 and 16.5 per cent. of copper; for the fineness of the French standard silver is 0.900 in the five-franc piece, while an inferior alloy of 0.835 is used for the lower denominations. The single franc-piece, composed of the latter alloy, is still made to weigh five grammes, the weight originally chosen for the franc as the unit of the monetary scale when the fineness of the coin was 0.900. It has now become a token, like the British shilling, of which the nominal value exceeds the metallic value. One pound troy of British standard silver is coined into 66

shillings, of which the metal is worth from 60*s*. to 62*s*. according to the market price of silver. The standard fineness of this silver is 0.925, three alloy in 40. British silver coins are a legal tender in payments to the amount of 40*s*. only.

## Chloride of Silver.

*Second Synthetical Reaction.*—To the product of the above reaction add water and hydrochloric acid or a soluble chloride; white chloride of silver (AgCl) is precipitated, copper still remaining in solution. Collect the precipitate on a filter, and wash with water; it is pure chloride of silver.

*Note.*—The nitrates of silver and copper may also be separated by evaporating the solution of the metals in nitric acid to dryness, and gently heating the residue, when the nitrate of copper is decomposed, but the nitrate of silver unaffected. The latter may be dissolved from the residual oxide of copper by water.

Chloride of silver may be obtained in crystals by evaporation of its solution in ammonia.

## Pure Silver.

*Third Synthetical Reaction.*—Place the chloride of silver of the previous reaction in a dish, wet it with dilute sulphuric acid, and float a piece of sheet zinc on the mixture; metallic silver is precipitated, and after about one day wholly removed from solution. Collect the precipitate on a filter and wash with water; it is pure metallic silver, and is readily fusible into a single button.

*Note.*—Any considerable quantity of chloride of silver may also be reduced to the metallic state by fusion, in a crucible, with about half its weight of carbonate of sodium.

## Pure Nitrate of Silver.

*Fourth Synthetical Reaction.*—Dissolve the pure silver of the previous reaction in nitric acid (3 of silver require about 2 or 2½ of strong acid diluted with 5 of water), and remove excess of acid by evaporating the solution to dryness, slightly heating the residue; the product is pure nitrate of silver. Dissolve by heating with a small quantity of water; on the solution cooling, or on evaporation, colorless tabular crystals of nitrate of silver are obtained.

| $3Ag_2$ | + | $8HNO_3$ | = | $2NO$ | + | $6AgNO_3$ | + | $4H_2O$ |
|---|---|---|---|---|---|---|---|---|
| Silver. | | Nitric acid. | | Nitric oxide. | | Nitrate of silver. | | Water. |

*Notes.*—The solution of pure or refined silver (*Argentum Purificatum*, B. P., *Argentum* U. S. P.) in nitric acid, evaporation, and

crystallization constitute the official process for the preparation of the nitrate (*Argenti Nitras*, B. P. and U. S. P.). The salt fused, and poured into proper moulds, yields the white cylindrical sticks or rods (*Argenti Nitras Fusa*, U. S. P.) commonly termed *caustic* (from *καίω*, *kaio*, I burn), or *lunar caustic*. (The alchemists called silver *Diana* or *Luna*, from its supposed mysterious connection with the moon.) The specimen of nitrate of silver obtained in the above reaction, dissolved in water, will be found useful as an analytical reagent. Nitrate of silver is soluble in rectified spirit; but after a time reaction and decomposition occur.

Silver salts are decomposed when in contact with organic matter, especially in the presence of light or heat, the metal itself being liberated, or a black insoluble compound formed. Hence the value of the nitrate in the manufacture of indelible ink for marking linen; hence, too, the reason of the practice of rendering silver solutions clear by subsidence and decantation, rather than by filtration through paper; and hence the cause of those cases of actual combustion which have been known to occur in preparing pills containing oxide of silver and essential oil or other organic matter. Linen marked with such ink should not be cleansed by aid of bleaching liquor, as the marked parts are then apt to be rapidly oxidized into perfectly rotten matter, holes resulting. Paul says the reaction is as follows: $Ag_2O + CaCl_2O_2 = 2AgCl + CaO + O_2$.

## Oxide of Silver.

*Fifth Synthetical Reaction.*—To a few drops of solution of nitrate of silver add solution of potash or soda or lime-water; an olive-brown precipitate of oxide of silver ($Ag_2O$) occurs. The washed and dried oxide, like most silver compounds, is decomposed by heat with production of metal.

The *Argenti Oxidum*, B. P. and U. S. P., is thus made, lime-water (solution of potash, U. S. P.) being the precipitant employed, soda and potash not being so readily removed by washing. Three and a half pints of lime-water will decompose half an ounce of nitrate of silver.

| $2AgNO_3$ | + | $Ca2HO$ | = | $Ag_2O$ | + | $Ca2NO_3$ | + | $H_2O$ |
|---|---|---|---|---|---|---|---|---|
| Nitrate of silver. | | Hydrate of calcium. | | Oxide of silver. | | Nitrate of calcium. | | Water. |

Methods of forming several other salts of silver are incidentally mentioned in the following analytical paragraphs.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—To a solution of a silver salt add hydrochloric acid or other soluble chloride; a white curdy precipitate of chloride of silver falls. Add nitric acid, and boil; the precipitate does not dissolve. Pour off the acid and add solution of ammonia; the precipitate dissolves. Neutralize

the ammoniacal solution by an acid; the chloride of silver is re-precipitated.

This is the most characteristic test for silver. The precipitated chloride is also soluble in solutions of hyposulphite of sodium or cyanide of potassium—facts of considerable importance in photographic operations.

*Other analytical reagents* than the above are occasionally useful.——Sulphuretted hydrogen, or sulphydrate of ammonium, gives a black precipitate, sulphide of silver ($Ag_2S$), insoluble in alkalies.——Solutions of potash or soda give a brown precipitate, oxide of silver ($Ag_2O$), converted into a fulminating compound by prolonged contact with ammonia.——Phosphate of sodium gives a pale yellow precipitate, phosphate of silver ($Ag_3PO_4$), soluble in nitric acid and in ammonia.——Arseniate of ammonium gives a chocolate-colored precipitate, arseniate of silver ($Ag_3AsO_4$), already noticed in connection with arsenic acid.——Iodide or bromide of potassium gives a yellowish-white precipitate, iodide or bromide of silver (AgI or AgBr), insoluble in acids and only slightly soluble in ammonia.——Cyanide of potassium gives a white precipitate, cyanide of silver (AgCy), soluble in excess, sparingly soluble in ammonia, insoluble in dilute nitric acid, soluble in boiling concentrated nitric acid. *Argenti cyanidum*, U. S. P., is made by distilling a mixture of ferrocyanide of potassium and diluted sulphuric acid, and passing the resulting hydrocyanic acid into a solution of nitrate of silver: $HCy + AgNO_3 = AgCy + HNO_3$ (the precipitate is well washed and dried).——Yellow chromate of potassium ($K_2CrO_4$) gives a red precipitate, chromate of silver ($Ag_2CrO_4$).——Red chromate of potassium also gives a red precipitate, acid chromate of silver ($Ag_2CrO_4,CrO_3$).——Many organic acids afford insoluble salts of silver.——Several metals displace silver from solution, mercury forming in this way a crystalline compound known as the silver tree, or *Arbor Dianæ*.——In the blowpipe flame, silver salts, placed on charcoal with a little carbonate of sodium, yield bright globules of metal accompanied by no incrustation as in the corresponding reaction with lead salts; the experiment may be performed with the nitrate, which first melts and then, like all nitrates, deflagrates, yielding a white metallic coating of silver which slowly aggregates to a button.

*Antidotes.*—Solution of common salt, sal-ammoniac, or any other inert chloride should obviously be administered where large doses of nitrate of silver have been swallowed. A quantity of sea-water or brine would convert the silver into insoluble chloride, and at the same time produce vomiting.

## QUESTIONS AND EXERCISES.

332. By what process is silver obtained from argentiferous galena?

333. What weight of English silver coin will yield one pound of pure nitrate of silver?

334. How may the metal be recovered from an impure mixture of silver salts?

335. Give a diagram showing the formation of nitrate of silver from the metal.

336. Describe the reaction of lime-water and nitrate of silver.

337. Mention the chief test for silver, and the precautions to be observed in order that silver salts may be distinguished from those of lead and mercury.

338. Name the antidote for silver.

---

### DIRECTIONS FOR APPLYING SOME OF THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE OF THE METALS, COPPER, MERCURY (EITHER AS MERCUROUS OR MERCURIC SALT), LEAD, SILVER.

Add hydrochloric acid:—

Silver is indicated by a white curdy precipitate, soluble in ammonia.

Mercurous salts also by a white precipitate, turned black by ammonia.

Lead by a white precipitate, insoluble in ammonia. Confirm by boiling another portion of the hydrochloric precipitate in water: it dissolves.

If hydrochloric acid gives no precipitate, silver and mercurous salts are absent. Lead can only be present in very small quantity. Mercuric salts may be present. Copper may be present. Divide the liquid into three portions, and apply a direct test for each metal as follows:—

Lead is best detected by the sulphuric test; the tube being set aside for a time if the precipitate does not appear at once.

Mercury is best detected by the copper test. If present, here it occurs as mercuric salt.

Copper betrays itself by the blue color of the liquid under examination. Confirm by the ammonia test.

If the above reactions are not thoroughly conclusive, confirmatory evidence should be obtained by the application of some of the other reagents for copper, mercury, lead, or silver.

TABLE OF SHORT DIRECTIONS FOR APPLYING SOME OF THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF **ANY OR ALL** OF THE METALS COPPER, MERCURY (EITHER MERCUROUS OR MERCURIC SALT, OR BOTH), LEAD, SILVER.

Add hydrochloric acid, filter, and wash the precipitate with a small quantity of cold water.

<table>
<tr><td colspan="2">Ppt.<br>Pb Hg(ous) Ag.<br>Wash with boiling water.</td><td rowspan="2">Filtrate<br>Cu Hg(ic) Pb.<br>Divide into three portions.<br>Test for<br>Cu by AmHO; blue sol.<br>Hg (mercuric) by Cu; globules.<br>Pb by $H_2SO_4$; white ppt.*</td></tr>
<tr><td colspan="2">Ppt.<br>Hg(ous) Ag.<br>Add AmHO.</td><td>Filtrate<br>Pb.<br>Add $H_2SO_4$, white ppt.*</td></tr>
<tr><td>Precipitate<br>Hg<br>(mercurous)<br>—black.</td><td>Filtrate<br>Ag.<br>Add $HNO_3$<br>white ppt.</td><td></td></tr>
</table>

* Liquids containing only a small quantity of lead do not readily yield suphate of lead on the addition of sulphuric acid. Before lead can be said to be absent, therefore, the liquid should be evaporated to dryness with one drop of sulphuric acid, and the residue digested in water; any sulphate of lead then remains as a heavy white insoluble powder.

OUTLINE OF THE PRECEDING TABLES.

| HCl | $H_2S$ | | AmHS | $Am_2CO_3$ | $Am_2HAsO_4$ | |
|---|---|---|---|---|---|---|
| Hg (as mercurous salt) | Cu | Insoluble in AmHS. | Zn | Ba | Mg | K |
| Pb (partially) | Hg (as mercuric salt) | | Al | Ca | | Na |
| Ag | Pb (entirely) | | Fe | | | Am |
| | As | Soluble in AmHS. | | | | |
| | Sb | | | | | |

The practical student should examine solutions containing the above metals until he is able to analyze with facility and accuracy. In this way he will best perceive the peculiarities of each element and their general relations to each other. As the rarer metals are not included here, the tables are not complete analytical schemes; only general memoranda respecting them will therefore now be given.

## Memoranda relating to the General Analytical Table (page 223).

*The group-tests* adopted in the Table are, obviously, hydrochloric acid, sulphuretted hydrogen, sulphydrate of ammonium, carbonate of ammonium, and arseniate of ammonium. If a group-test produces no precipitate, it is self-evident that there can be no member of the group present. At first, therefore, add only a small quantity of a group test, and if it produces no effect add no more; for it is not advisable to overload a solution with useless reagents; substances expected to come down as precipitates are not unfrequently held in the liquid by excess of acid, alkali, or strong aqueous solution of some group-reagent, thoughtlessly added. Indeed, experienced manipulators not unfrequently make preliminary trials with group-reagents on a few drops only of the liquid under examination; if a precipitate is produced, it is added to the bulk of the original liquid and the addition of the group-reagent continued; if a precipitate is not produced, the few drops are thrown away, and the unnecessary addition of a group-reagent thus avoided altogether, an advantage fully making up for the extra trouble of making

a preliminary trial.——While shunning excess, however, care must be taken to avoid deficiency; a substance only partially removed from solution through the addition of an insufficient amount of a reagent will appear where not expected, be consequently mistaken for something else, and cause much trouble; this will not occur if the appearance, odor, or reaction of the liquid on test-paper be duly observed. It is also a good plan, when a group-reagent has produced a precipitate and the latter has been filtered out, to add a little more of the reagent to the clear filtrate; if more precipitate is produced, an insufficient amount of the group-test was introduced in the first instance; but the error is corrected by simply refiltering; if no precipitate occurs, the mind is satisfied and the way cleared for further operations.

*Group-precipitates*, or any precipitates still requiring examination, should, as a rule, be well washed before further testing; this is to remove the aqueous solution of other substances adhering to the precipitate (the mother-liquor, as it is termed), so that subsequent reaction may take place fairly between the reagent used and the precipitate only.——A precipitate is sometimes in so fine a state of division as to retard filtration by clogging the pores of the paper, or even to pass through the filter altogether; in these cases the mixture may be warmed or boiled (or a fresh quantity of the original solution may be warmed *before* the group-test is added), which usually causes aggregation of the particles of a precipitate, and hence facilitates the passage of liquids.

*Division of work.*—It is immaterial whether a solution be first divided into group-precipitates or each precipitate be examined as soon as produced; if the former method be adopted, confusion will be avoided by labelling or marking the funnels or papers holding the precipitate "the HCl ppt.," "the $H_2S$ ppt.," and so on.

*The colors and general appearance* of the various sulphides and hydrates precipitated should be borne in mind, as the absence of other bodies, as well as the presence of those thrown down, is often at once thus indicated.

*Application of confirmatory tests* must be frequent.

*Results of analysis* should be recorded neatly in a memorandum-book.

*The various reactions* which occur in an analysis have already come before the reader in going through the tests for the individual metals or in other analytical operations; it is unnecessary, therefore, again to draw out equations or dia-

grams. But the reactions should be thought over, and, if not perfectly clear to the mind, be written out again and again till thoroughly understood.

---

## QUESTIONS AND EXERCISES.

339. Give processes for the qualitative analysis of liquids containing the following substances:—

*a*. Antimony and Mercurous salt.
*b*. Lead and Calcium.
*c*. Silver and Mercurous salt.
*d*. Lead and Mercuric salt.
*e*. Copper and Arsenicum.
*f*. Arsenicum and Antimony.
*g*. Aluminium and Zinc.
*h*. Iron and Copper.
*i*. Magnesium, Calcium, and Potassium.
*j*. Silver, Antimony, Zinc, Barium, and Ammonium.

340. Enumerate the so-called group-tests.

341. Give a general sketch of the method of analyzing a solution suspected to contain two or more salts of common metals.

342. Classify the common metals according to their analytical relations.

---

# METALS OF MINOR PHARMACEUTICAL IMPORTANCE.

Thus far has been considered, somewhat in detail, the chemistry of the common metals, salts of which are frequently used in medicine or in testing medical substances. These are—

| | | | | |
|---|---|---|---|---|
| Potassium, | Barium, | Zinc, | Arsenicum, | Mercury, |
| Sodium, | Calcium, | Aluminium, | Antimony, | Lead, |
| Ammonium (?), | Magnesium, | Iron, | Copper, | Silver. |

Of the remaining metals, eight are mentioned in the British Pharmacopœia, namely:—

| | | | |
|---|---|---|---|
| Lithium, | Chromium, | Gold, | Cadmium, |
| Manganese, | Tin, | Platinum, | Bismuth. |

Compounds of three more are sufficiently common to occasionally come under notice—

Strontium, Cobalt, Nickel.

These eleven metals of minor pharmaceutical interest may be shortly studied, a few only of the reactions of each (just those mentioned in the following pages) being performed. When all have

been thus treated, their respective positions in the analytical groups will be indicated and a tabular scheme by which an analysis of a solution containing any metal may be effected. Thus, step by step, we may learn how to analyze almost any substance that may occur, and know to what extent the presence of a rarer will interfere with the ordinary tests for a common element; additional illustrations of the working of chemical laws will be acquired, and the store of chemical and pharmaceutical facts increased. The opportunity thus afforded for improvement in habits of neatness in manipulation, precision, and classification furnishes another and no mean reason why such experiments should be prosecuted, the direct value of which may not be considerable to medical and pharmaceutical learners.

## LITHIUM.

Symbol L. Atomic weight 7.

Lithium is widely distributed in nature, but usually in minute proportions compared with other elements. A trace of it may be found in most soils and waters, a Cornish spring containing even considerable quantities as chloride.

One salt used in medicine is the *Citrate* ($L_3C_6H_5O_7$) (*Lithii Citras*, U. S. P.), occurring in white deliquescent crystals or powder prepared by dissolving 50 grains of the *Carbonate* ($L_2CO_3$) and 100 of citric acid (50 to 95 if both are quite pure) in 1 ounce of water, evaporating to a low bulk, and setting aside in a dry place to crystallize, or at once evaporating to dryness and powdering the residue. The crystals have the formula $L_3C_6H_5O_7,4H_2O$; dried at 212° F., $L_3C_6H_5O_7,H_2O$ (Umney).

$$\underset{\text{Carbonate of lithium.}}{3L_2CO_3} + \underset{\text{Citric acid.}}{2H_3C_6H_5O_7} = \underset{\text{Citrate of lithium.}}{2L_3C_6H_5O_7} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

The carbonate *Lithii Carbonas*, U. S. P.) is a white granular powder obtained from the minerals which contain lithium; namely, lepidolite (from λεπίς, *lepis*, a scale, and λίθος, *lithos*, a stone; it has a scaly appearance), triphane (from τρεῖς, *treis*, three, and φαίνω, *phainō*, I shine), or spodumene (from σποδόω, *spodōō*, I reduce to ashes, in allusion to its exfoliation in the blowpipe-flame), and petalite (from πέταλον, *petalon*, a leaf; its character is leafy and laminated). Each contains silicate of aluminium, with fluoride of potassium and lithium in the case of lepidolite, and silicate of sodium and lithium in the others. *Liquor Lithiæ Effervescens*, B. P., is a solution of 10 grains of carbonate of lithium in 1 pint of water charged with 7 times its volume of carbonic acid gas and kept in ordinary aerated-water bottles. "Half a pint, evaporated to dryness, yields 5 grains of a white solid residue, answering to the tests for carbonate of lithium. . . . Ten grains of the latter salt neutralized with sulphuric acid, and afterwards heated to redness, leave 14.86 grains of dry sulphate of lithium, which, when redissolved in distilled water, yields no precipitate with oxalate of ammonium or solution of lime," indicating absence of salts of calcium and aluminium. Citrate of

lithium should yield by incineration 52.8 per cent. of white carbonate of lithium.

*Urate of lithium** is more soluble than urate of sodium; hence lithium preparations are administered to gouty patients in the hope that urate of sodium, with which such systems are loaded, may be converted into urate of lithium and removed.

*In chemical position* lithium stands between the alkaline and the alkaline-earth metals, its hydrate, carbonate, and phosphate being slightly soluble in water. The double chloride of platinum and lithium also is soluble in water. Its atom is univalent, L'.

*Analytical Reaction.*—Moisten the end of a platinum wire with solution of a minute particle of solid lithium salt, and introduce it into the flame of a Bunsen burner or other slightly colored flame (spirit-lamp or blowpipe-flame); a magnificent crimson tinge is imparted.

The light emitted by ignited lithium vapor is of a purer scarlet than that given by strontium, the next element. When the flames are examined by spectral analysis (physically analyzed by a prism), the red rays are, in the case of strontium, found to be associated with blue and yellow, neither of which is present in the lithium light.

## STRONIUM.

Symbol Sr. Atomic weight 87.5.

*Source.*—Strontium is not widely distributed in nature; but the carbonate ($SrCO_3$) known as *strontianite*, and the sulphate ($SrSO_4$) known as *celestine* (from *cœlum*, the sky, in allusion to its occasional bluish color), are by no means rare minerals.

*Salts of strontium* are not employed in medicine. They are chiefly used by firework manufacturers in preparing red fire. The color they impart to flame is a beautiful crimson—ignited strontium vapor emitting red rays, as already explained. Nitrate of strontium ($Sr2NO_3$) is best for pyrotechnic compositions, its oxygen enabling it to burn freely when mixed with charcoal, sulphur, etc. It, or any salts, may be obtained by dissolving the carbonate in the appropriate acid, or by igniting the cheaper sulphate with coal, whereby sulphide (SrS) is produced, and dissolving this in acid.

*The position of strontium* among the chemical elements is between barium and calcium; its sulphate is very sparingly soluble in water. Its atom, like those of barium and calcium, is bivalent (Sr'').

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a solution of a strontium salt ($Sr2NO_3$ or $SrCl_2$) add carbonate of ammonium; a white precipitate of carbonate of strontium ($SrCO_3$) falls.

---

* Urates will be considered subsequently in connection with uric acid.

*Second Analytical Reaction.*—To a solution of a strontium salt add sulphuric acid previously so diluted that it will not precipitate calcium salts or an equally dilute solution of any sulphate (*e. g.*, that of calcium itself); a white precipitate of sulphate of strontium ($SrSO_4$) falls. The formation of this precipitate is promoted by stirring and by setting the liquid aside for some time.

Barium is precipitated immediately under similar circumstances.

*Third Analytical Reaction.*—To a dilute solution of a strontium salt add yellow chromate of potassium; no precipitate falls.

Barium may be separated from strontium by chromate of potassium, that reagent at once precipitating barium from aqueous or acetic solutions.

*Fourth Analytical Reaction.*—Insert a fragment of a strontium salt in the blowpipe-flame, or other equally colorless flame, or hold the end of a platinum wire dipped into a strontium solution in the flame; a crimson color is imparted.

*Other Analytical Reactions.* — Alkali-metal phosphates, arseniates, and oxalates give white insoluble precipitates with strontium as with barium and calcium.——Strontium, like calcium, but unlike barium, is not precipitated by hydrofluosilicic acid.

CERIUM. Ce. At. wt. 22.—This element occurs in the mineral cerite (a silicate of iron, calcium, and the three rare metals, cerium, lanthanum, and didymium); also occasionally as impure fluoride, carbonate, and phosphate. The oxalate of cerium, a white granular powder, is the only official salt; it may be obtained from cerite by boiling the powdered mineral in strong hydrochloric acid for several hours, evaporating, diluting, and filtering to separate silica; adding ammonia to precipitate hydrates of all the metals except calcium; filtering off, washing, redissolving in hydrochloric acid, and adding oxalic acid to precipitate oxalate of cerium. The preparation will still contain oxalates of lanthanum and didymium; it is therefore strongly calcined, the resulting oxides of lanthanum and didymium dissolved out to some extent by boiling with a concentrated solution of chloride of ammonium, the residual oxide of cerium dissolved in boiling hydrochloric acid, and oxalate of ammonium added to precipitate oxalate of cerium ($Ce''C_2O_4,3H_2O$).

Oxalate of cerium (*Cerii Oxalas*, B. P. and U. S. P.) is decomposed at a dull red heat, 48 per cent. of a yellow, or, more generally, a salmon-colored mixture of oxides remaining; usually the didymium present gives the ignited residue a reddish or reddish-brown color; it is then soluble in boiling hydrochloric acid (without effervescence; indicating, indirectly, absence of earthy and other carbonates or oxalates), and the solution gives, with excess of a saturated solution

of sulphate of potassium, a crystalline precipitate of double sulphate of cerium and potassium. Alumina mixed with oxalate of cerium may be detected by boiling with solution of potash, filtering, and adding excess of solution of chloride of ammonium, when a white flocculent precipitate of hydrate of aluminium will be obtained. The oxalic radical is recognized by neutralizing the potash solution by acetic acid and adding chloride of calcium; white oxalate of calcium is then precipitated; this precipitate, though insoluble in acetic, should be wholly dissolved by hydrochloric acid.

According to H. G. Greenish, most samples of oxalate of cerium have as impurities traces of lead, iron, and magnesium.

## MANGANESE.

Symbol Mn. Atomic weight 55.

*Source.*—Manganese is a constituent of many minerals, and as black oxide ($MnO_2$) (*Manganesii Oxidum Nigrum*, B. P. and U. S. P.) or *pyrolusite* (from πῦρ, *pur*, fire, and λύσις, *lusis*, a loosing or resolving, in allusion to the readiness with which it is split up by heat into a lower oxide and oxygen) occurs frequently in abundance in the southwest of England, Aberdeenshire, and most of the countries of Europe. It is met with as a steel-gray mass of prismatic crystals, or in black shapeless lumps.

*The chemical position* of manganese is close to iron and three other metals still to be considered—cobalt, nickel, and chromium. Its atom apparently has sexivalent affinities, as seen in manganate of potassium ($K_2MnO_4$); but commonly it is quadrivalent ($Mn^{IV}$) or bivalent ($Mn''$).

*Uses.*—Metallic manganese is only used in alloy with iron in the manufacture of some varieties of steel. The black oxide is an important agent in the production of chlorine, the preparation of green and red disinfecting manganates, purple glass, and black glazes for earthenware.

### *Reactions having both Synthetical and Analytical Interest.*

*First Reaction.*—Boil a few grains of black oxide of manganese with some drops of hydrochloric acid until chlorine ceases to be evolved; add water, and filter; the filtrate is a solution of manganous chloride ($MnCl_2$).

$$MnO_2 + 4HCl = MnCl_2 + 2H_2O + Cl_2.$$

This is the reaction commonly applied in the preparation of chlorine gas. It is also a ready method of preparing a manganous salt for analytical experiments. Coupled with the application of reagents to the filtrate, the reaction is that by which a black powder or mineral would be recognized as black oxide of manganese. Black oxide of manganese dissolves in cold hydrochloric acid, forming a dark brown solution of a higher chloride or chlorides, $MnCl_3$, $Mn_2Cl_7$, or, possibly, $MnCl_4$.

*Second Reaction.*—Heat a particle of a manganese compound with a grain or two of carbonate and hydrate of potassium and a fragment of nitrite or chlorate of potassium on platinum foil in the blowpipe-flame; a green mass containing *manganate of potassium* ($K_2MnO_4$) results. Boil the foil in a little water; the green manganate dissolves and soon changes to solution of the purple *permanganate of potassium* ($K_2Mn_2O_8$).

This is a delicate analytical test for manganese.

The reaction is similar to that by which permanganate of potassium (*Potassii Permanganas*, U. S. P.) is directed to be prepared for use in volumetric analysis. *Liquor Potassii Permanganatis*, U. S. P., is a solution of 80 grains of permanganate of potassium in 1 pint of distilled water. Equations showing the exact action which occurs in making the salt according to the process of the British Pharmacopœia have already been given in connection with the compounds of potassium (*vide* p. 76). The proportions of ingredients and details of the operation are as follows:—

Reduce 3½ parts (for experiment each "part" may be ½th oz.) of chlorate of potassium to fine powder, and mix it with 4 of black oxide of manganese; put the mixture into a porcelain basin, and add to it 5 parts of solid caustic potash, previously dissolved in 4 parts of water. Evaporate to dryness, stirring diligently to prevent spirting. Pulverize the mass, put it into a covered Hessian or Cornish crucible, and expose it to a dull red heat (not higher) for an hour (20 or 30 minutes for quantities of 1 or 2 ozs.), or till it has assumed the condition of a semifused mass. Allow to cool, pulverize, and boil with about 30 parts of water. Let the insoluble matter subside, decant the fluid, boil again with about 10 parts of water, again decant, neutralize the united liquors accurately with diluted sulphuric acid (or, better, carbonic acid gas), and evaporate till a pellicle forms. Set aside to cool and crystallize. Drain the crystalline mass, boil it in 6 parts of water, and strain through a funnel the throat of which is lightly obstructed by a little asbestos or gun-cotton. Let the fluid cool and crystallize, drain the dark purple slender prismatic crystals, and dry them by placing under a bell jar over a vessel containing sulphuric acid.

Instead of converting the manganate into permanganate by ebullition, by which one-third of the manganate is lost, Städeler recommends chlorine to be passed through the cold solution until the green color is entirely changed to purple.

*Solutions of the manganates* of potassium and sodium are in common use as disinfectants under the name of Condy's fluid. They act by oxidizing organic matter, the manganic or permanganic radical being reduced to black manganic oxide, or even a lower oxide. The reason for using asbestos instead of paper in filtering the solutions will now be understood.

The changes in color which the green mass of the above process undergoes when dropped into warm water procured for it the old name of *mineral chameleon*.

*Third Reaction.*—Make a borax bead by heating a fragment of the salt on the looped end of a platinum wire in the blow-pipe-flame until a clear transparent globule is obtained. Place on the bead a minute portion of a manganese compound, or touch it with a drop of solution. Again fuse the borax, using the point of the flame; a bead of a violet or amethystine tint is produced.

This is a good analytical reaction. It has also synthetical interest, illustrating the use of black oxide of manganese in producing common purple-tinted glass.

Expose the bead to the reducing part of the flame, the part nearer to the blowpipe, where there are highly heated hydrocarbon gases greedy of oxygen; the color disappears.

This is owing to the reduction of the manganic compound to a manganous condition, in which it no longer possesses peculiar coloring-power. This action also illustrates the use of black oxide of manganese in glass-manufacture. Glass when first made is usually of a green tint, owing to the presence of ferrous impurities; the addition of manganic oxide to the materials converts the ferrous into ferric compounds, which have comparatively little colorific power, it itself being thereby reduced to manganous oxide, which also gives but little color. If excess of manganic oxide be added, a purple tint is produced.

### *Reactions having Analytical Interest (Tests).*

*Fourth Reaction.*—Through a solution of a manganous salt acidified by hydrochloric acid pass sulphuretted hydrogen; no decomposition occurs. Add ammonia; the sulphydrate of ammonium thus formed causes the precipitation of a yellowish-pink or flesh-tinted precipitate of manganous sulphide (MnS) in a hydrous state.

This reaction is characteristic, sulphide of manganese being the only flesh-colored sulphide known. The salt used may be the manganous chloride obtained in the first reaction; but such crude solutions usually give a black precipitate with sulphydrate of ammonium, owing to the presence of iron. The latter element may be precipitated, however, on adding excess of ammonia (and rapidly filtering, oxygen will be absorbed and most of the manganese also precipitated) or on boiling the manganous solution with a very little carbonate of sodium, which attacks the ferric salt in preference to the manganous. *Pure* manganous chloride may be similarly obtained on boiling the impure solution with manganous carbonate; the latter decomposes the ferric chloride with production of ferric hydrate and more manganous chloride, and evolution of carbonic acid gas.

To the recently precipitated manganous sulphide add acetic acid; it is dissolved.

This solubility enables manganese to be separated from nickel, cobalt, and zinc, whose sulphides are insoluble in weak acetic acid. To express the fact in another way—manganese is not precipitated by sulphuretted hydrogen from a solution containing free acetic acid only.

*Fifth Reaction.*—To solution of manganous salt add ammonia; a white precipitate of manganous hydrate (Mn2HO) falls. Add excess of ammonia; some of the precipitate is dissolved and may be detected in the quickly filtered solution by sulphydrate of ammonium. But both precipitate and solution rapidly absorb oxygen, the manganese passing into a more highly oxidized condition in which it is insoluble in ammonia.

The fixed alkalies give a similar precipitate *insoluble* in excess. The precipitate rapidly absorbs oxygen, becomes brown, and gradually passes into a higher oxide.

*Sixth Reaction.*—Heat a little black oxide of manganese in a test-tube with sulphuric acid; oxygen is evolved and sulphate of manganese formed (*Manganesii Sulphas*, U. S. P.); add water, boil, filter, evaporate, and set aside to crystallize. Larger quantities are made in a similar manner.

Sulphate of manganese ($MnSO_4,5H_2O$) occurs in colorless, or pale rose-colored, transparent crystals, which, when deposited from a solution at a temperature between 68° and 86°, have the form of right rhombic prisms, and contain four molecules of water. This salt is very soluble in water. The solution is not colored by tincture of nutgall a black (a black shows iron). but affords with caustic alkalies a white precipitate (Mn2HO), which, by exposure to the air, soon absorbs oxygen, and becomes brown. Sulphydrate of ammonium throws down a flesh-colored precipitate (MnS), and ferrocyanide of potassium a white one ($Mn_2Fcy$).

*Many other reactions* occur between manganese salts and various reagents, but are of no particular synthetical or analytical interest.

A good method proposed by Crum, for detecting minute quantities of manganese, consists in adding dilute nitric acid and either red lead or the puce-colored oxide or peroxide of lead to the solution, and then boiling; a red tint, said to be due to permanganic acid, is imparted to the liquid. If chlorides are present the manganese, etc., should be separated by potash or soda, the precipitate be well washed, dissolved in nitric acid, and then the oxide be added.

## COBALT.

Symbol Co. Atomic weight 58.8.

*Source.*—Cobalt occurs sparingly in nature as the arsenide ($CoAs_2$), or *tin-white cobalt*, and occasionally as a double arsenide and sulphide ($CoAs_2,CoS_2$), or *cobalt-glance* (from *glanz*, brightness, in allusion to its lustre).

*Uses.*—Its chief use is in the manufacture of blue glass, the color of which is due to a compound of cobalt. Cobalt is also the coloring constituent of *smalt* (from *smelt*, a corruption of *melt*), a finely-ground sort of glass, used as a blue pigment by paper-stainers and others, and employed also by laundresses to neutralize the yellowish appearance of washed linen.

*The salts of cobalt* may be obtained from the oxide (CoO), and the oxide from *zaffre*, a mixture of sand and roasted ore.

*Quantivalence.*—The atom of cobalt often exhibits quadrivalent affinities, but still more often exerts only bivalent powers (Co″). Cobalt has analytical relations with zinc, nickel, and manganese, and may be regarded as a member of the iron group.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—Pass sulphuretted hydrogen through a solution of a salt of cobalt—the chloride ($CoCl_2$) or nitrate (Co2NO$_3$) for example; no decomposition occurs. Add ammonia; the sulphydrate of ammonium thus formed causes the precipitation of black sulphide of cobalt (CoS).

The moist precipitate slowly absorbs oxygen from the air, becoming converted into sulphate of cobalt ($CoSO_4$).

*Second Analytical Reaction.*—Add ammonia gradually to a cobalt solution; a blue precipitate of impure hydrate of cobalt (Co2HO) falls. Add excess of ammonia; the precipitate is dissolved, yielding a liquid somewhat more reddish-brown than the original solution.

A similar precipitate is given by the fixed alkalies, *insoluble* in excess.

*Third Analytical Reaction.*—Make a borax bead by heating a fragment of the salt on the looped end of a platinum wire in a blowpipe-flame until a clear transparent globule is obtained. Place on the bead a minute portion of a cobalt compound, or touch it with a drop of solution. Again fuse the borax; a blue bead results.

This is a delicate test for cobalt. From what has previously been said, it will be seen that this experiment has also considerable synthetical interest.

*Fourth Analytical Reaction.*—To a solution of a salt of cobalt add two or three drops of hydrochloric acid, then excess

of solution of cyanide of potassium, and boil for ten minutes; oxygen is absorbed, and cobalticyanide of potassium ($K_6Co_2Cy_{12}$) formed. Add hydrochloric acid, and boil the mixture (in a fume-cupboard, to avoid inhalation of any hydrocyanic acid); the excess of cyanide of potassium is thus decomposed, but the cobalticyanide is unaffected. Now add excess of solution of potash; the cobalticyanide of potassium is decomposed, but the cobalt remains dissolved in the alkaline liquid.

Nickel under similar circumstances is precipitated, the reaction thus affording means of separating these closely allied metals from each other.

*Other reactions* between a cobalt solution and different reagents may be performed, and various precipitates obtained; but these have no special analytical interest.

*Invisible Ink.*—Many salts of cobalt containing water of crystallization are light red, the anhydrous more or less blue. Prove this by writing some words on paper with a solution of chloride of cobalt sufficiently dilute for the characters to be invisible when dry; hold the sheet before a fire or over a flame; the letters at once become visible, distinct, and of a blue color. Breathe on the words, or set the sheet aside for a while; the characters are once more invisible, owing to absorption of moisture. Hence solution of chloride of cobalt forms one of the so-called *sympathetic inks.*

## NICKEL.

Symbol Ni. Atomic weight 58.8.

Nickel is, chemically, closely allied to cobalt, the ores of the two metals being commonly associated in nature. Indeed, it is from *speiss*, an arsenio-sulphide of nickel obtained in the manufacture of smalt, a pigment of cobalt already mentioned, that most of the nickel met with in commerce is obtained. It is much used in the preparation of the white alloy known as German or nickel silver.

*Quantivalence.*—Nickel exerts bivalent activity (Ni″) in its ordinary compounds. Its salts and their solutions are usually green. They are chiefly made, directly or indirectly, from the metal itself.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—Pass sulphuretted hydrogen through a solution of a salt of nickel—chloride ($NiCl_2$), nitrate ($Ni2NO_3$), or sulphate ($NiSO_4$); no decomposition occurs. Add ammonia; the sulphydrate of ammonium thus formed causes the precipitation of black sulphide of nickel (NiS).

*Note.*—When sulphate of nickel is precipitated by the direct addition of the common yellow solution of sulphydrate of ammonium, which always contains free sulphur, there is much difficulty in filtering the mixture, owing to the slight solubility of sulphide of nickel in the reagent and the formation of some sulphate of nickel ($NiSO_4$), oxygen being absorbed from the air by the sulphide. This may be avoided by warming the mixture and using freshly-made sulphydrate of ammonium, in which the sulphide of nickel is insoluble; or, where practicable, the salt of nickel may be precipitated from an ammoniacal solution by sulphuretted hydrogen.

*Second Analytical Reaction.*—Add ammonia drop by drop to a nickel solution; a pale-green precipitate of hydrate of nickel (Ni2HO) falls. Add excess of ammonia; the precipitate dissolves, yielding a bluish rather than the original green-colored solution.

A similar precipitate is given by the fixed alkalies, *insoluble* in excess.

*Third Analytical Reaction.*—Nickel salts color a borax bead, when hot, a reddish-yellow tint; the reaction is not very serviceable analytically.

*Fourth Analytical Reaction.*—To a solution of a salt of nickel add solution of cyanide of potassium; cyanide of nickel ($NiCy_2$) is precipitated. Add excess of solution of cyanide of potassium; the precipitate is dissolved with formation of double cyanide of nickel and potassium ($NiCy_2$,2KCy). Next add hydrochloric acid, and boil the mixture (in a fume-cupboard), adding a little hydrochloric acid from time to time until all smell of hydrocyanic acid has disappeared. Lastly, add excess of solution of potash; hydrate of nickel is precipitated.

### *Qualitative Separation of Cobalt and Nickel.*

The foregoing reaction serves for the separation of nickel from cobalt. On adding excess of hydrochloric acid to a solution containing the two metals, together with cyanide of potassium, a precipitate of cyanide of nickel and cobalticyanide of nickel occurs. By ebullition with excess of hydrochloric acid the cyanide of nickel is decomposed, chloride of nickel going into solution. On then adding excess of potash, hydrate of nickel is precipitated. The cobalticyanide of nickel is not decomposed by the acid; but is by the alkali, its cobalt going into solution and its nickel remaining insoluble as hydrate.

After filtering off the nickel, cobalt is detected in the filtrate by evaporating to dryness and testing the residue with borax in the blowpipe-flame.

*Another process.*—To any solution containing cobalt and nickel add excess of ammonia, and then enough solution of ferridcyanide of potassium to dissolve any precipitate. If

cobalt be present a reddish-brown color is produced in the liquid, due to the formation and solution of ferridcyanide of cobalt. Now add solution of ferrocyanide of potassium; a whitish precipitate of ferrocyanide of nickel appears, either at once, or, if the liquid is very weak, on *partially* neutralizing the free ammonia by hydrochloric acid. The ammonia must not be entirely neutralized by acid, nor enough acid added to decompose the reddish ferridcyanide of cobalt. If these precautions be not observed, a greenish-white precipitate of ferrocyanide of cobalt may be formed, and be mistaken for the ferrocyanide of nickel.

The value of this method (Skey and Davies) turns on the facts that ferridcyanide of nickel is not a colored body, while ferridcyanide of cobalt is reddish-brown, and that ferridcyanogen has apparently, *in ammoniacal solution*, greater affinity for cobalt than for nickel, while ferrocyanogen has, apparently, greater affinity for nickel than for cobalt. The formulæ of these so-called ferrocyanides and ferridcyanides of cobalt and nickel have not been definitely ascertained.

*Other reactions* between a nickel solution and various reagents give, in many cases, insoluble precipitates which, from their green color, are occasionally useful in distinguishing nickel from allied elements.

## CHROMIUM.

Symbol Cr. Atomic weight 52.5.

*Source.*—The chief ore of chromium is chrome ironstone, a mixture of the oxides of the metals ($FeO,Cr_2O_3$), occurring chiefly in the United States and Sweden. In constitution it seems to resemble magnetic iron ore ($FeO,Fe_2O_3$).

*Preparation of Red Chromate of Potassium.*—On roasting the powdered ore with carbonate of potassium and nitre, yellow chromate of potassium ($K_2CrO_4$) is obtained; the mass, treated with acid, yields red or bichromate ($K_2CrO_4,CrO_3$) (*Potassii Bichromas*, U. S. P.); from this salt other chromates are prepared, and by reduction, as presently explained, the salts of chromium itself. The yellow and orange chromates of lead are largely used as pigments.

*Note on Constitution.*—Red chromate of potassium is a somewhat abnormal salt, containing, probably, neutral chromate associated with chromic anhydride. The value of chromates as chemical reagents is alluded to in connection with chromate of barium (p. 103). Heated strongly in a crucible, red chromate of potassium splits up into yellow chromate, glistening oxide of chromium, and oxygen.

*Quantivalence.*—Chromium stands in close chemical relation to iron, aluminium, and manganese. Its atom is sexivalent if the formula of the fluoride ($CrF_6$) be correct. Like iron and aluminium, it

is trivalent, as seen in chromic chloride ($Cr_2Cl_6$), but sometimes exerts only bivalent activity, as in chromous chloride ($CrCl_2$).

*Passage of Chromium from the Acidulous to the Basylous Side of Salts.*—Through an acidified solution of red chromate of potassium pass sulphuretted hydrogen; sulphur is deposited, and a green salt of chromium remains in solution, chloride ($Cr_2Cl_6$) if hydrochloric acid be used, and sulphate ($Cr_2 3SO_4$) if sulphuric be the acid employed. Boil the liquid to expel excess of sulphuretted hydrogen, filter, and reserve the solution for subsequent experiments. (For an equation of this reaction, see p. 239.)

Alcohol, sugar, or almost any substance which is tolerably liable to oxidation, will answer as well as sulphuretted hydrogen.

*Sulphate of chromium* ($Cr_2 3SO_4$), like sulphate of aluminium ($Al_2 3SO_4$), unites with alkali-metal sulphates to form *alums*, which resemble common alum both in crystalline form and, as far as we know, in internal structure: they are of purple color.

## Reactions.

*Chromium* as *Chromic Acid*, or other *Chromate*.—This is the state in which chromium will usually be met with, the most common salt being the red chromate or bichromate of potassium. Mix four volumes of a cold, saturated aqueous solution of red chromate of potassium with five of oil of vitriol; on cooling, *chromic anhydride* ($CrO_3$), *Acidum Chromicum*, U. S. P., separates in crimson needles. After well draining, the crystals may be freed from adhering sulphuric acid by washing once or twice with nitric acid: the latter may be removed by passing dried and slightly warmed air through a tube containing the crystals. In contact with moisture chromic anhydride takes up water and forms solution of true chromic acid ($H_2CrO_4$). Chromic anhydride is a powerfully corrosive oxidizing agent. It melts between 356° and 374° F.

The oxygen in chromic acid and other chromates, and in manganates, permanganates, black oxide of manganese, and puce-colored oxide of lead is in a physically different state from that in peroxide of hydrogen, peroxide of barium, and similar compounds. On bringing chromic acid or the above acidified solution of red chromate of potassium into contact with solution of peroxide of hydrogen, a strong effervescence of oxygen ensues. According to Schönbein and Brodie the oxygen of chromic acid is in the negative or ozonic state, while that of peroxide of hydrogen is in the positive or so-called antozonic condition. Both are equally active, but neutralize each other, forming neutral or ordinary oxygen.

In the analytical examination of solutions containing chromates, the chromium will always come out in the state of green chromic hydrate along with ferric hydrate and alumina, the prior treatment by sulphuretted hydrogen reducing the chromium in the molecule to the lower state, thus:—

$$K_2CrO_4,CrO_3 + 8HCl + 3H_2S = Cr_2Cl_6 + 2KCl + 7H_2O + S_3.$$

Chromium having been found in a solution, its condition as chromate may be ascertained by applying to the original solution salts of barium, mercury, lead, and silver. (See the various paragraphs relating to those metals.)

| | | | |
|---|---|---|---|
| $Ba2NO_3$ | gives yellow | $BaCrO_4$ | with chromates. |
| $Hg_2NO_3$ | " red | $Hg_2CrO_4$ | " |
| $AgNO_3$ | " " | $Ag_2CrO_4$ | " |
| " | " " | $Ag_2CrO_4,CrO_3$ | with bichromates. |
| $Pb2C_2H_3O_2$ | " yellow | $PbCrO_4$ | with both. |

Nitrate of barium does not completely precipitate bichromates, bichromate of barium being soluble in water; the chromate of barium is insoluble in water or acetic acid, but soluble in hydrochloric or nitric acid. Mercurous nitrate does not wholly precipitate bichromates: mercuric nitrate or chloride only partially precipitates chromates, and does not precipitate bichromates. The mercurous chromate is insoluble, or nearly so, in diluted nitric acid. The silver chromates are soluble in acids and alkalies. Acetate of lead precipitates chromates and bichromates, acetic acid being set free in the latter case.

*A delicate reaction for dry chromates* will be found in the formation of *chlorochromic anhydride* ($CrO_2Cl_2$). A small portion of the chromate is placed in a test-tube with a fragment of dry chloride of sodium and a drop or two of oil of vitriol, and the mixture heated; red irritating fumes of chlorochromic *anhydride* are evolved, and condense in dark red drops on the side of the tube.

Large quantities of pure distilled chlorochromic anhydride are obtained by the same reaction, the operation being conducted in a retort, with thoroughly dry materials, for the compound is decomposed by water. It may be regarded as chromic anhydride in which an atom of oxygen is displaced by an equivalent quantity (two atoms) of chlorine. It is not used in medicine, but is of interest to the chemical student as being an illustration of a large class of similar bodies—*chloro-acidulous* or *chloro-anhydro compounds*. The reaction is also occasionally serviceable for the detection of chlorides.

### *Analytical Reactions of Chromium Salts (Tests).*

*First Analytical Reaction.*—To solution of a salt of chromium (chloride, sulphate, or chrome alum) add sulphydrate of ammonium; a bulky green precipitate of chromic hydrate ($Cr_2 6HO$), containing a large quantity of water (7 molecules, $7H_2O$), is precipitated.

$$Cr_2Cl_6 + 6AmHS + 6H_2O = Cr_2 6HO + 6AmCl + 6H_2S.$$

*Second Analytical Reaction.*—To solution of a chromium salt add ammonia; chromic hydrate is precipitated, insoluble in excess.

*Third Analytical Reaction.*—To solution of a chromium salt add solution of potash or soda drop by drop; chromic hydrate is precipitated. Add excess of the fixed alkali; the precipitate is dissolved. Boil well the solution; the chromic hydrate is reprecipitated.

*Iron, Chromium, and Aluminium Salts*, chemically so alike, may be separated by this reaction. Ferric hydrate is insoluble in solutions of the fixed alkalies, cold or hot; chromium hydrate soluble in cold but not in hot; hydrate of aluminium in both. To a solution containing all three metals, therefore, add potash or soda, stir, and filter; the iron is thrown out: boil the filtrate, and filter; the chromium is thrown out: neutralize the filtrate by acid, and then add ammonia; the aluminium is thrown out. Note, however, that ferric hydrate will prevent hydrate of chromium being dissolved by potash or soda if the ferric hydrate is in considerable excess. Before concluding that chromium is entirely absent, the 4th reaction should be performed. The hydrates of iron, chromium, and aluminium are insoluble in ammonia, and may therefore be easily separated from the hydrates of the somewhat analogous metals zinc, cobalt, nickel, and manganese.

*Fourth Analytical Reaction.*—Add a salt of chromium (either of the above precipitates of chromic oxide or the dry residue of the evaporation of a few drops of a solution of a chromium salt) to a few grains of nitre and carbonate of sodium on platinum foil, and fuse the mixture in the blowpipe-flame; a yellow mass of chromate of potassium and sodium ($KNaCrO_4$) is formed. Dissolve the mass in water, add acetic acid to decompose excess of carbonate, and apply the reagents for chromates.

This is a delicate and useful reaction if carefully performed.

## TIN.

Symbol Sn. Atomic weight 118.

*Source.*—The chief ore of tin is stannic oxide ($SnO_2$), occurring in veins under the name of *tinstone*, or in alluvial deposits as *stream-tin*. The principal mines are those of Cornwall.

*Preparation.*—The metal is obtained by reducing the roasted and washed ore by charcoal or anthracite* coal at a high temperature, and is purified by slowly heating, when the pure tin, fusing first, is run off, a somewhat less fusible alloy of tin with small quantities of arsenicum, copper, iron, or lead remaining. The latter is known as *block tin;* the former heated till brittle and then hammered or let fall from a height splits into prismatic fragments, resembling starch or columnar basalt, and is named *dropped* or *grain tin*. Good tin emits a crackling noise in bending, termed the cry of tin, caused by the friction of its crystalline particles on each other.

*Uses.*—Tin is an important constituent of such alloys as pewter, Britannia metal, solder, speculum-metal, bell-metal, gun-metal, and bronze. It is very ductile, and may be rolled into plates or leaves, known as *tin foil*, varying from $\frac{1}{250}$ to $\frac{1}{1000}$ of an inch in thickness. Common tin foil, however, usually contains a large proportion of lead. The reflecting surface of most *looking-glasses* is an amalgam of tin and mercury, produced by carefully sliding a plate of glass over a sheet of tin foil on which mercury has been rubbed, and then excess of mercury poured. *Pins* are made of brass wire, on which tin is deposited. *Tin plate*, of which common utensils are made, is iron alloyed with tin by dipping the cleansed sheet into melted tin. *Tin tacks* are in reality tinned iron tacks; a tin nail would be too soft to drive into wood. Tin may be granulated by melting and triturating briskly in a hot mortar, by shaking melted tin in a box on the inner sides of which chalk has been rubbed, or, in thin little bells or corrugated fragments (Granulated Tin, B. P.), by melting in a ladle and, as soon as fluid, pouring from the height of a few feet into water. Powdered tin has been used medicinally as a mechanical irritant to promote expulsion of worms. The hair of the pods of *Kiwach* or *Cowhage* (Hindustani) (*Mucuna pruriens*, P. I.) is almost the only medicine (excluding diluents and dentifrices) which acts in such a directly mechanical manner.

*The chemical position of tin* among the metals is close to that of arsenicum and antimony. Its atom is quadrivalent and bivalent. The two classes of salts are termed stannic and stannous respectively. They are all made directly or indirectly from the metal itself.

---

* *Anthracite* (from ἄνθραξ, *anthrax*, a burning coal) or *stone coal* differs from the ordinary *bituminous* or *caking coal*, in containing less volatile matter, and, therefore, in burning without flame. It gives a higher temperature, and from its non-caking properties is, in furnace operations, more manageable than bituminous coal.

## Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

### (*a*) *Synthetical Reactions.*

### Chloride of Tin. Stannous Chloride.

*First Synthetical Reaction.*—Warm a fragment of tin with hydrochlorous acid; hydrogen escapes and solution of stannous chloride ($SnCl_2$) is formed. It may be retained for future experiments.

One ounce of tin dissolved in three fluidounces of hydrochloric acid and one of water, and the resulting solution diluted to five fluidounces, constitutes the "Solution of the Chloride of Tin," B. P.

*Solid Stannous Chloride.*—By evaporation of the above solution stannous chloride is obtainable in crystals ($SnCl_2,2H_2O$). It is a powerful reducing agent, even a dilute solution precipitating gold, silver, and mercury from their solutions, converting ferric and cupric into ferrous and cuprous salts, and partially deoxidizing arsenic, manganic, and chromic acids. It absorbs oxygen from the air, and is decomposed when added to a large quantity of water unless some acid be present. It is used as a mordant in dyeing and calico-printing.

### Perchloride of Tin. Stannic Chloride.

*Second Synthetical Reaction.*—Through a portion of the solution of the stannous chloride of the previous reaction pass chlorine gas; solution of stannic chloride ($SnCl_4$) is formed. Or add hydrochloric acid to the stannous solution, boil, and slowly drop in nitric acid until no more fumes are evolved; again stannic chloride results. Reserve the solutions for subsequent experiments.

### Stannic Oxide, or Anhydride, and Stannates.

*Third Synthetical Reaction.*—Boil a fragment of tin with nitric acid, evaporate to dryness, and strongly calcine the residue; light buff-tinted stannic anhydride ($SnO_2$) is produced. Heat the stannic anhydride with excess of solid caustic potash or soda; stannate of the alkali metal ($K_2SnO_3$ or $Na_2SnO_3$) results. Dissolve the stannate in water, and add hydrochloric acid; white, gelatinous *stannic acid* ($H_2SnO_3$) is precipitated. Stannic acid is also obtained on adding an alkali to solution of stannic chloride; it is soluble in excess of acid or alkali.

The product of the action of nitric acid on tin is also an acid, but from its insolubility in hydrochloric and other acids, is different from

ordinary stannic acid. It is termed *metastannic acid* (from μετὰ, *meta*, beyond), and probably has a composition expressed by the formula $H_{10}Sn_5O_{15}$. (*Vide* Index, "Isomerism.") It is also produced on gently heating stannic acid:—

$$\underset{\text{Stannic acid.}}{5H_2SnO_3} = \underset{\text{Metastannic acid.}}{H_{10}Sn_5O_{15}}$$

Metastannates have the general formula $M_2H_8Sn_5O_{15}$. Both acids yield buff-colored stannic oxide or anhydride ($SnO_2$) when strongly heated. The latter is employed in polishing plate under the name of *putty powder*. *Stannate of sodium* ($Na_2SnO_3,4H_2O$) is used as a mordant by dyers and calico-printers under the name of *tin prepare-liquor*.

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*Stannous or stannic salts.*—Heat any solid tin compound with a mixture of cyanide of potassium and carbonate of sodium on charcoal by the inner flame of the blowpipe. Globules of tin separate, having, when cut by a knife, characteristic brightness and hardness.

#### STANNOUS SALTS.

*First Analytical Reaction.*—Through a solution of a stannous salt (stannous chloride, for example, see previous page) pass sulphuretted hydrogen gas; brown stannous sulphide (SnS) is precipitated. Pour off the supernatant liquid, add ammonia to the moist precipitate (to neutralize acid), and lastly yellow sulphydrate of ammonium; the precipitate is dissolved.

Aqueous solution of sulphydrate of ammonium becomes yellow when a day or two old, and then contains excess of sulphur, that element having become displaced by oxygen absorbed from the air; hence, in the above reaction, the stannous sulphide (SnS), in dissolving, becomes stannic sulphide ($SnS_2$); for the latter is precipitated on decomposing the alkaline liquid by an acid.

*Second Analytical Reaction.*—To solution of a stannous salt add solution of potash or soda; white stannous hydrate falls (Sn2HO). Add excess of the alkali; the precipitate dissolves. Boil the solution; some of the tin is precipitated as black stannous oxide (SnO).

Ammonia gives a similar precipitate, insoluble in excess. The alkaline carbonates do the same, carbonic acid gas escaping.

### STANNIC SALTS.

*Third Analytical Reaction.*—Through a solution of a stannic salt (stannic chloride, for example, see page 242) pass sulphuretted hydrogen gas; yellow stannic sulphide ($SnS_2$) is precipitated. Pour off the supernatant liquid, and to the moist precipitate add ammonia (to neutralize acid) and then sulphydrate of ammonium; the precipitate dissolves.

*Note.*—In precipitating stannic sulphide the presence of too much hydrochloric acid must be avoided; the formation of the precipitate is also facilitated if the solution be warmed. Stannic sulphide, like the sulphide of arsenicum and antimony, dissolves in a solution of alkaline sulphide or sulphydrate, with formation of definite crystallizable sulphostannates ($M'_2SnS_3$).

*Anhydrous stannic sulphide*, prepared by sublimation, has a yellow or orange lustrous appearance, and is used by decorators as *bronzing-powder*. It is sometimes termed *mosaic gold.*

*Fourth Analytical Reaction.*—To solution of a stannic salt add potash or soda; white stannic acid falls ($H_2SnO_3$). Add excess of the alkali; the precipitates dissolves. Boil the mixture; no reprecipitation occurs—a fact enabling stannic to be distinguished from stannous salts.

Ammonia gives a similar precipitate, soluble, but not readily, in excess. The fixed alkali-metal carbonates do the same, carbonic acid gas escaping; after a time the stannic salt is again deposited, probably as stannate of the alkali metal. Carbonate of ammonium, and acid carbonates of alkali metals give a precipitate of stannic acid insoluble in excess.

*Antidotes.*—In cases of poisoning by tin salts (dyer's tin liquor *e. g.*), solution of carbonate of ammonium should be given. White of egg is also said to form an insoluble precipitate with compounds of tin. Vomiting should be speedily induced, and the stomach-pump quickly applied.

## GOLD.

Symbol Au. Atomic weight 196.7.

*Source.*—Gold occurs in the free state in nature, occasionally in nodules or *nuggets*, but commonly in a finer state of division termed *gold dust.*

*Preparation.*—Gold is separated from the sand, crushed quartz, or other earthy matter with which it may be associated, by agitation with water, when the gold, from its relatively greater specific gravity, falls to the bottom of the vessels first, the lighter mineral matter being allowed to run off with the water. From this rich sand the gold is dissolved out by mercury, the amalgam filtered, and afterwards distilled, when the mercury volatilizes and gold remains. The amalgamation may be much facilitated by the use of a small proportion of sodium, as described under silver (page 216).

*Pure gold* is too soft for general use as a circulating medium. *Gold coin* is an alloy of copper and gold, that of Great Britain containing 1 of the former to 11 of the latter, or 8⅓ per cent. of copper, that of France, Germany, and the United States about 10 per cent. *Jeweller's gold* varies in quality, every 24 parts containing 18, 15, 12, or 9 parts of gold, the alloys being technically termed 18, 15, 12, or 9 *carat fine*. Articles made of the better qualities are usually stamped by authority. Trinkets of inferior intrinsic worth are commonly thinly coated with pure gold by electro-deposition or otherwise. *Gold leaf* is nearly pure gold passed between rollers till it is about $\frac{1}{1000}$ of an inch in thickness and then hammered between sheets of animal membrane, termed gold-beater's skin and calf-skin vellum, till it is $\frac{1}{250000}$ or $\frac{1}{280000}$ of an inch in thickness. It may even be hammered till 280,000 leaves would be required to form a pile an inch thick.

*Gold Coinage.*—The weight of gold is expressed in Great Britain in ounces troy and decimal parts of an ounce, and the metal is always taken to be of standard fineness (11 gold and 1 alloy) unless otherwise described. The degree of fineness of gold, as ascertained by assay, is expressed decimally, fine pure gold ("gold free from metallic impurities," B. P.) being taken as unity, or 1.000. Thus gold of British standard is said to be 0.9166 fine, of French standard 0.900 fine. The legal weight of the sovereign is 0.2568 ounce of standard gold, or 123.274 grains. The weight came from one pound of standard gold (5760 grains) being coined into 44½ guineas. Gold coins are legal tender to any amount, provided that the weight of each sovereign does not fall below 122.5 grains, or in the case of a half sovereign 61.125 grains; these are the "least current" weights of the coins.

*Note.*—In chemical analysis gold comes out among the sulphides of the metals precipitated by sulphuretted hydrogen; and of those sulphides, it, like the sulphides of tin, antimony, and arsenicum, is soluble in sulphydrate of ammonium.

*Quantivalence.*—Gold is trivalent (Au‴), but in some compounds univalent (Au′).

## Reactions.

*Synthetical Reaction.*—Place a fragment of gold (*e. g.*, gold leaf) in ten or twenty drops of aqua regia (a mixture of three parts of nitric and four or five of hydrochloric acid), and set the test-tube aside in a warm place; solution of perchloride of gold or auric chloride ($AuCl_3$) results. When the metal is dissolved, evaporate nearly to dryness to remove most of the excess of fluid, dilute with water, and retain the solution for subsequent experiments. Sixty grains of gold treated thus, and the resulting chloride dissolved in five ounces of distilled water, constitute "Solution of Chloride of Gold," B. P.

$$Au_2 + 2HNO_3 + 6HCl = 2AuCl_3 + 2NO + 4H_2O.$$

This reaction has analytical interest also; for in examining a substance suspected to be or contain metallic gold, solution would have to be effected in the above way before reagents could be applied. Gold is insoluble in hydrochloric, nitric, and the weaker acids.

### *Analytical Reactions (Tests.)*

*First Analytical Reaction.*—Through a few drops of solution of an auric salt (the chloride, $AuCl_3$, is the only convenient one) pass sulphuretted hydrogen; brown auric sulphide ($Au_2S_3$) is precipitated. Filter, wash, and add yellow sulphydrate of ammonium solution; the precipitate dissolves.

*Second Analytical Reaction.*—To solution of a salt of gold add ferrous chloride or sulphate, and set the tube aside; metallic gold is precipitated, a ferric salt remaining in solution.

This is a convenient way of preparing pure gold, or *fine gold* as it is termed, or of working up the gold residues of laboratory operations. The precipitate, after boiling with hydrochloric acid, washing and drying, may be obtained in a button by mixing with an equal weight of borax or acid sulphate of potassium and fusing in a good furnace.

*Third Analytical Reaction.*—Add a few drops of dilute solutions of stannous and stannic chloride to a considerable quantity of distilled water; pour the liquid, a small quantity at a time, into a dilute solution of auric chloride ($AuCl_3$), well stirring; the mixture assumes a purple tint, and flocks of a precipitate, known as the *Purple of Cassius* (from the name of the discoverer, M. Cassius) are produced.

The same compound is formed on immersing a piece of tin foil in solution of auric chloride; it is said to be a mixture of auric, aurous, stannic, and stannous oxides. It is the coloring agent in the finer varieties of ruby glass.

## PLATINUM.

Symbol Pt. Atomic weight 198.

*Source.*—Platinum, like gold, usually occurs in nature in the free state, the chief sources of supply being Mexico, Brazil, and Siberia. It is separated from the alluvial soil by washing.

*Uses.*—The chief use of platinum is in the construction of foil, wire, crucibles, spatulas, capsules, evaporating-dishes, and stills, for the use of the chemical analyst or manufacturer. It is tolerably hard, fusible with very great difficulty, not dissolved by hydrochloric, nitric, or sulphuric acid, and only slightly affected by alkaline substances. It is attacked by aqua regia with production of perchloride of platinum or platinic chloride ($PtCl_4$). It forms fusible alloys with lead and other metals, and with phosphorus a phosphide, which easily

melts. Neither of these substances, therefore, nor mixtures which may yield a metal, should be heated in platinum vessels.

*The chemical position of platinum* among the elements is close to that of gold. Its atom is quadrivalent in some compounds, in others apparently bivalent (Pt''). The higher salts are termed *platinic*, the lower *platinous*.

The specific gravity of platinum is 21.5; and that of iridium, an allied metal, 22.4.

## Reactions.

### Perchloride of Platinum. Platinic Chloride.

*Synthetical Reaction.*—Place a fragment of platinum in a little aqua regia and set the vessel aside in a warm place, adding more acid from time to time if necessary; solution of perchloride of platinum ($PtCl_4$) results. Evaporate the solution to remove excess of acid, and complete the desiccation over a water-bath. Dissolve the residue in water and retain the solution for subsequent experiments, and as a reagent for the precipitation of salts of potassium and ammonium.

A quarter of an ounce of platinum treated in the above manner, and the resulting chloride dissolved in five ounces of water, constitutes "Solution of Perchloride of Platinum," B. P.

This reaction has analytical interest also; for in examining a substance suspected to be or to contain metallic platinum, solution would have to be thus effected before reagents could be applied.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—Through a few drops of a solution of a platinic salt ($PtCl_4$ is the only convenient one) to which an equal quantity of solution of chloride of sodium has been added, pass sulphuretted hydrogen; dark brown platinic sulphide ($PtS_2$) is precipitated. Filter, wash, and add sulphydrate of ammonium; the precipitate dissolves.

If chloride of sodium be not present in the above reaction, the precipitated sulphide will contain platinous chloride, and may detonate when heated.

*Second Analytical Reaction.*—Add excess of solution of carbonate of sodium and some sugar to solution of perchloride of platinum and boil; a precipitate of metallic platinum falls.

*Platinum Black* (B. P.) is the name of this precipitate. It possesses in a high degree a quality common to many substances, but largely possessed by platinum, namely, that of absorbing or occluding gases. In its ordinary state, after well washing and drying, it absorbs from the air and retains many times its bulk of oxygen. A

drop of ether or alcohol placed on it is rapidly oxidized, the platinum becoming hot. This action may be prettily shown by pouring a few drops of ether into a beaker (one having portions of the top and sides broken off answers best), loosely covering the vessel with a card, and suspending within the beaker a platinum wire, one end being attached to the card by passing through its centre, the other terminating in a short coil or helix near the surface of the ether; on now warming the helix in a flame and then rapidly introducing it into the beaker, it will become red-hot and continue to glow so long as there is ether in the vessel. In this experiment real combustion goes on between the ether vapor and the concentrated oxygen of the air, the products of the oxidation revealing themselves by their odor.

*Third Analytical Reaction.*—To solution of perchloride of platinum add solution of chloride of ammonium; a yellow granular precipitate of double chloride of platinum and ammonium ($PtCl_4,2AmCl$) falls. When slowly formed in dilute solutions, the precipitate is obtained in minute orange prisms.

Chloride of potassium (KCl) gives a similar precipitate ($PtCl_4$-$2KCl$). Platinic chloride having been stated to be a test for potassium and ammonium salts, the reader is prepared to find that potassium and ammonium salts are tests for platinic salts. The double sodium compound ($PtCl_4,2NaCl$) is soluble in water.

Collect the precipitate, dry, and heat in a small crucible; it is decomposed, and metal, in the finely divided state of *spongy platinum*, remains.

$$3(PtCl_4,2NH_4Cl) = Pt_3 + 2NH_4Cl + 16HCl + 2N_2.$$

Heat decomposes the potassium salt into $Pt + 2KCl + Cl_4$, the chlorine escaping and the chloride of potassium remaining with the platinum.

*In working up the platinum residues of laboratory operations*, the mixture should be dried, burnt, boiled successively with hydrochloric acid, water, nitric acid, water, then dissolved in aqua regia, excess of acid removed by evaporation, chloride of ammonium added, the precipitate washed with water, dried, ignited, and the resulting spongy platinum retained or converted into perchloride for use as a reagent for alkali metals. It is by this process that the native platinum is treated to free it from the rare metals palladium, rhodium, osmium, ruthenium, and iridium. The spongy platinum is converted into the massive condition by a refinement on the blacksmith's process of *welding* (German *wellen*, to join), or by fusing in a flame of pure oxygen and hydrogen gases, the oxyhydrogen blowpipe.

*Occlusion by Spongy Platinum.* — Spongy platinum has great power of occlusion. A small piece held in a jet of hydrogen causes ignition of the gas, owing to the close approximation of particles of oxygen (from the air) and hydrogen. Döbereiner's lamp is con-

structed on this principle—the apparatus being essentially a vessel in which hydrogen is generated by the action of diluted sulphuric acid on zinc, and a cage for holding the spongy platinum.

## CADMIUM.

Symbol Cd. Atomic weight 112.

In most of its chemical relations cadmium (*Cadmium*, U. S. P.) resembles zinc. In nature it occurs chiefly as an occasional constituent of the ores of that metal. In distilling zinc containing cadmium, the latter, being the more volatile, passes over first. In analytical operations, cadmium, unlike zinc, comes down among the metals precipitated by sulphuretted hydrogen; that is, its sulphide is insoluble in dilute hydrochloric acid, while sulphide of zinc is soluble. It is a white malleable metal nearly as volatile as mercury. Sp. gr. 8.7.

Beyond the occasional employment of the sulphide as a pigment (*jaune brillant*), and the iodide in photography and medicine, cadmium and its salts are but little used. The atom of cadmium is bivalent (Cd″).

### Reactions.

### Iodide of Cadmium.

*First Synthetical Reaction.*—Digest together in a flask metallic cadmium, water, and iodine until the color of the iodine disappears; solution of iodide of cadmium (*Cadmii Iodidum*, B. P.) ($CdI_2$) remains. Pearly micaceous crystals may be obtained on evaporating the solution.

This is the process alluded to in the British Pharmacopœia. The compound is used in medicine in the form of ointment, *Unguentum Cadmii Iodidi*, B. P. The salt is also employed, with other iodides, in iodizing collodion for photographic purposes. It melts when heated, and is soluble in water or spirit, the solution reddening litmus paper.

### Sulphate of Cadmium.

*Second Synthetical Reaction.*—Dissolve cadmium in nitric acid; pour the resulting solution of nitrate of cadmium ($Cd2NO_3$) into a solution of carbonate of sodium; dissolve the precipitate of carbonate of cadmium ($CdCO_3$) in dilute sulphuric acid, separate and crystallize. Sulphate of cadmium ($CdSO_4$) is a white crystalline salt soluble in water (U. S. P.).

*First Analytical Reaction.*—Through solution of a cadmium salt ($CdI_2$ or $CdCl_2$) pass sulphuretted hydrogen; a yellow precipitate of sulphide of cadmium (CdS) falls, resembling in appearance arsenious, arsenic, and stannic sulphides. Add sulphydrate of ammonium; the precipitate, unlike the sulphides just mentioned, does not dissolve.

Sulphides of cadmium and copper may be separated by solution of cyanide of potassium, in which sulphide of copper is soluble and sulphide of cadmium insoluble.

*Second Analytical Reaction.*—To a cadmium solution add solution of potash; white hydrate of cadmium (Cd2HO) is precipitated, insoluble in excess of the potash.

Hydrate of zinc (Zn2HO), precipitated under similar circumstances, is soluble in solution of potash; the filtrate from the hydrate of cadmium may therefore be tested for any zinc occurring as an impurity by applying the appropriate reagent—sulphydrate of ammonium.

*Before the blowpipe-flame*, on charcoal, cadmium salts give a brown deposit of oxide of cadmium (CdO).

## BISMUTH.

Symbol Bi. Atomic weight 208.

*Source.*—Bismuth occurs in the metallic state in nature. It is freed from adherent quartz, etc., by simply heating, when the metal melts, runs off, and is collected in appropriate vessels. It is also met with in combination with other elements. Bismuth is grayish-white, with a distinct pinkish tinge.

*Uses.*—Beyond the employment of some of its compounds in medicine, bismuth is but little used. Melted bismuth expands considerably on solidifying, and hence is valuable in taking sharp impressions of dies. It is a constituent of some kinds of type-metal and of pewter-solder.

*The position of bismuth among the metals* is close to that of arsenicum and antimony. Its atom is rarely quinquivalent ($Bi^{V}$), but in most compounds trivalent ($Bi'''$).

### Reactions having (*a*) Synthetical and (*b*) Analytical Interest.

#### (*a*) *Reactions having Synthetical Interest.*

#### Nitrate of Bismuth.

*First Synthetical Reaction.*—To a few drops of nitric acid and an equal quantity of water in a test-tube, add a little powdered bismuth, heating the mixture if necessary; nitric oxide (NO) escapes, and solution of *nitrate of bismuth* ($Bi3NO_3$) results.

$$\underset{\text{Bismuth.}}{Bi_2} + \underset{\text{Nitric acid.}}{8HNO_3} = \underset{\text{Nitrate of bismuth.}}{2(Bi3NO_3)} + \underset{\text{Nitric oxide.}}{2NO} + \underset{\text{Water.}}{4H_2O}$$

The solution evaporated gives crystals ($Bi3NO_3,5H_2O$), any arsenicum which the bismuth might contain remaining in the mother-liquor. Native bismuth (*Bismuthum*, B. P. and U. S. P.) commonly contains arsenicum, most of which is removed by roasting or by fusing two or three times with a tenth of its weight of nitre (*Bismuthum Purificatum*, B. P.), or, finally, by converting the metal into oxynitrate, as described in the next reaction, and reducing this with charcoal at a high temperature.

To make *nitrate* of bismuth and other salts on a larger scale, 2 ounces of the metal, in small fragments, are gradually added to a mixture of 4 fluidounces of nitric acid and 3 of water, and, when effervescence (due to escape of nitric oxide) has ceased, the mixture is heated for ten minutes, poured off from any insoluble matter, evaporated to 2 fluidounces to remove excess of acid, and then either set aside for *crystals* to form, or poured into a half gallon of water to form the *oxynitrate* of bismuth, or into a solution of 6 ounces of carbonate of ammonium in a quart of water to form the *oxycarbonate* as described in the following reactions. The precipitates should be washed with cold water and dried at a temperature not exceeding 150° F. Exposed in the moist state to 212° for any length of time, they undergo slight decomposition.

## Subnitrate or Oxynitrate of Bismuth.

*Second Synthetical Reaction.*—Pour some of the above solution of nitrate into a considerable quantity of water; decomposition occurs and oxynitrate of bismuth ($BiONO_3$) in a hydrous state ($BiONO_3,H_2O$) (*Bismuthi Subnitras*, B. P.) is precipitated;—

$$\underset{\text{Nitrate of bismuth.}}{Bi3NO_3} + \underset{\text{Water.}}{H_2O} = \underset{\text{Oxynitrate of bismuth.}}{BiONO_3} + \underset{\text{Nitric acid.}}{2HNO_3}$$

Filter, and test the filtrate for bismuth by adding excess of carbonate of sodium; a precipitate shows that some bismuth remains in solution. The following equation, therefore, probably more nearly represents the decomposition:—

$$\underset{\text{Nitrate of bismuth.}}{5(Bi3NO_3)} + \underset{\text{Water.}}{8H_2O} = \underset{\text{Oxynitrate of bismuth.}}{4(BiONO_3.H_2O)} + \underset{\text{Nitrate of bismuth in acid.}}{Bi3NO_3, 8HNO_3}$$

Decomposition of nitrate of bismuth by water is the process of the British Pharmacopœia for the preparation of oxynitrate or "subnitrate" of bismuth for use in medicine. For this purpose the original metal must contain no arsenicum. In manufacturing the compound, therefore, before pouring the solution of nitrate into water, the liquid should be tested for arsenicum by one of the hydrogen tests; if that element be present, the solution must be evaporated and only the deposited crystals be used in the preparation of the oxynitrate. For on pouring an arsenical solution of nitrate of bismuth into water, the arsenicum is not wholly removed in the super-

natant liquid, unless the oxynitrate be redissolved and reprecipitated several times, according to the amount of arsenicum present

In the United States Pharmacopœia the solution of nitrate (made from washed carbonate) is directed to be poured into dilute solution of ammonia. Such a product will probably contain more oxyhydrate than oxynitrate.

*Subnitrate of Bismuth* is sometimes administered in the form of a lozenge (*Trochisci Bismuthi*, B. P.). It is used as a cosmetic under the name of *Pearl-white* (*Blanc de Perle*).

*Oxysalts of Bismuth.*—It will be noticed that the formula for subnitrate of bismuth ($BiNO_4$) does not accord with that of other nitrates, the characteristic elements of which are $NO_3$. Analogy would seem to indicate, however, that the fourth atom of oxygen has different functions from the three in the $NO_3$; for on pouring solution of chloride of bismuth ($BiCl_3$) into water, oxychloride is produced (BiOCl) (a white powder used as a cosmetic, also in enamels, and in some varieties of sealing-wax). The bromide ($BiBr_3$) and iodide ($BiI_3$) similarly treated yield oxybromide (BiOBr) and oxyiodide (BiOI). The subnitrate ($BiNO_4$) is, therefore, probably an analogous compound, an oxynitrate ($BiONO_3$). The sulphate ($Bi3SO_4$) also decomposes when placed in water, giving what may be termed an oxysulphate ($Bi_2O_2SO_4$).

It is difficult to prove whether or not the water in the "subnitrate" or hydrous oxynitrate of bismuth ($BiONO_3,H_2O$) is an integral part of the salt. If it is, the compound is simply the hydrato-nitrate ($BiNO_32HO$) of bismuth.

## Oxide of Bismuth.

*Third Synthetical Reaction.*— Boil subnitrate of bismuth with solution of soda for a few minutes; it is converted into yellowish oxide of bismuth ($Bi_2O_3$) (*Bismuthi Oxidum*, B. P.).

$$\underset{\text{Oxynitrate of bismuth.}}{2BiONO_3} + \underset{\text{Hydrate of sodium.}}{2NaHO} = \underset{\text{Oxide of bismuth.}}{Bi_2O_3} + \underset{\text{Nitrate of sodium.}}{2NaNO_3} + \underset{\text{Water.}}{H_2O}$$

## Subcarbonate or Oxycarbonate of Bismuth.

*Fourth Synthetical Reaction.*— To solution of nitrate of bismuth add carbonate of ammonium; a white precipitate of hydrous oxycarbonate ($2Bi_2O_2CO_3,H_2O$) (*Bismuthi Carbonas*, B. P.) falls.

$$\underset{\text{Nitrate of bismuth.}}{4(Bi3NO_3)} + \underset{\text{"Carbonate" of ammonium.}}{3N_4H_{16}C_3O_9} = \underset{\text{Nitrate of ammonium.}}{12NH_4NO_3} + \underset{\text{Oxycarbonate of bismuth.}}{2Bi_2O_2CO_3} + \underset{\text{Carbonic acid gas.}}{7CO_2}$$

According to the United States Pharmacopœia the oxycarbonate (*Bismuthi Subcarbonas*) is made by precipitating with

carbonate of sodium a solution of the nitrate prepared from washed hydrate of bismuth.

This compound may be regarded as similar in constitution to the oxysalts just described. In $Bi_2CO_5$ one scarcely recognizes the characteristic elements of carbonates; but considering the preparation to be an oxycarbonate ($Bi_2O_2CO_3$) its relations to carbonates and oxides are evident. These subsalts may all be viewed as normal bismuth salts in which an atom of oxygen displaces an equivalent proportion of other acidulous atoms or radicals:—

| | | | |
|---|---|---|---|
| Chloride . . . . | $Bi3Cl$ | Oxychloride . . | $BiOCl$ |
| Bromide . . . . | $Bi3Br$ | Oxybromide . . | $BiOBr$ |
| Iodide . . . . . | $Bi3I$ | Oxyiodide . . | $BiOI$ |
| Nitrate . . . . . | $Bi3NO_3$ | Oxynitrate . . | $BiONO_3$ |
| Sulphate . . . . | $Bi_2 3SO_4$ | Oxysulphate . | $Bi_2O_2SO_4$ |
| Carbonate (unknown) | $Bi_2 3CO_3$ | Oxycarbonate . | $Bi_2O_2CO_3$ |

They may be viewed, in short, as salts in process of conversion to oxide; continue the substitution a little further, and each yields oxides of bismuth ($Bi_2O_3$). They have also been considered to be salts of a hypothetical univalent radical bismuthyl (BiO).

## Citrate of Bismuth.

*Fifth Synthetical Reaction.*—To solution of nitrate of bismuth add citric acid and then solution of ammonia until the precipitate at first formed is redissolved, and the liquid after shaking has a slight ammoniacal odor. The product contains citrate of bismuth ($BiC_6H_5O_7$) dissolved in solution of citrate and nitrate of ammonium. Made with definite quantities of ingredients and an amount of bismuth salt equivalent to the three grains of oxide ($Bi_2O_3$) in a fluidrachm, the solution forms the *Liquor Bismuthi et Ammoniæ Citratis*, B. P.

If the ordinary oxynitrate of bismuth be heated for a few moments with a concentrated solution of an equivalent of citric acid, the normal citrate is generated as a heavy crystalline powder, and the nitric acid entirely freed and easily washed away by decantation. Ten parts of the oxynitrate, seven of citric acid crystals, and thirty to forty of water are heated together for a few minutes, until a drop of the mixture forms a clear solution with ammonia water, the crystalline mass is then diluted with eight to ten times its volume of water, and set aside for a short time to let the citrate deposit; the clear liquid is then decanted. The crystalline sediment is then washed three or four times in a similar manner, drained, and dried either on a water-bath or by mere exposure. The yield is 13⅔ parts, showing that the salt is anhydrous, and that its formula is $BiC_6H_5O_7$ (Rother).

### (*b*) *Reactions having Analytical Interest* (*Tests*).

*First Analytical Reaction.*—Through solution of a bismuth salt (a slightly acid solution of nitrate, for example) pass sulphuretted hydrogen; a black precipitate of sulphide of bismuth ($Bi_2S_3$) falls. Add ammonia (to neutralize acid) and then sulphydrate of ammonium; the precipitate, unlike $As_2S_3$ and $Sb_2S_3$, is insoluble.

*Second Analytical Reaction.*—Concentrate almost any acid solution of a bismuth salt and pour into water; a white salt is precipitated.

This reaction is characteristic of bismuth salts; it has already been amply explained. The precipitate is distinguished from one formed by antimony under similar circumstances, by being insoluble in solution of tartaric acid.

*Third Analytical Reaction.*—To a solution of a bismuth salt add an alkali; hydrate of bismuth (Bi3HO) is precipitated insoluble in excess.

*Fourth Analytical Reaction.*—A small quantity of the following reagent, including both supernatant liquid and precitated scales, is transferred to a test-tube and gradually heated until solution takes place. Any liquid containing or supposed to contain bismuth is then added, and the whole allowed to cool. The separated scales will show a distinct change in color to dark orange or crimson according to the quantity of bismuth present.

The *test-reagent* may be prepared by adding to a boiling solution of acetate of lead (half a grain to the ounce) solution of iodide of potassium in considerable excess. The solution of the iodide of lead precipitated is assisted by a little acetic acid. On cooling, iodide of lead is deposited in the characteristic scales.

---

The reader is again advised to trace out the exact nature of each of the foregoing reactions, chiefly by aid of equations or diagrams.

---

## QUESTIONS AND EXERCISES.

343. Enumerate the fifteen metals, salts of which are frequently employed in pharmacy.

344. Mention the eleven rarer metals.

345 Name the sources and official compounds of lithium.

346. Give an equation explanatory of the formation of Citrate of Lithium.

347. What is the strength of *Liquor Lithiæ Effervescens?*

348. On what chemical hypothesis are lithium compounds administered to gouty patients?

349. Describe the relation of lithium to other metals.

350. What is the chief test for lithium?

351. Write a paragraph on strontium, its natural compounds, chemical relations, technical applications, and tests.

352. What are the formulæ and properties of oxalate of cerium?

353. Name the commonest ore of manganese, and give an equation descriptive of its reaction with hydrochloric acid.

354. Explain the formation of permanganate of potassium, employing diagrams or equations.

355. In what manner do the manganates of potassium act as disinfectants?

356. What are the chief tests for manganese?

357. What are the chief uses of the compounds of cobalt?

358. How are salts of cobalt analytically distinguished from those of nickel?

359. Mention an application of nickel in the arts.

360. What is the general color of nickel salts?

361. State the method of preparation of red chromate of potassium.

362. Give the formulæ of red and yellow chromates of potassium.

363. How is red chromate of potassium obtained?

364. Describe the action of sulphuretted hydrogen on acidified solutions of chromates.

365. What is the formula of chrome alum?

366. Mention the chief tests for the chromic radical, and for chromium.

367. How would you detect iron, chromium, and aluminium in a solution?

368. Define the terms tinstone, stream-tin, block-tin, grain-tin, tin-plate.

369. Describe the position occupied by tin in relation to other metals.

370. What is the difference between stannic acid and metastannic acid?

371. State the applications of tin in the arts.

372. Mention the chief tests for stannous and stannic salts.

373. Name the best antidote in cases of poisoning by tin solution.

374. How is gold dust separated from the earthy matter with which it is naturally associated?

375. How much pure gold is contained in English coin, and in jewellers' gold?

376. State the average thickness of gold leaf.

377. What is the weight of a sovereign?

378. Explain the term "fineness" as applied to gold.

379. What effect is produced on gold by hydrochloric, nitric, and nitrohydrochloric acids respectively?

380. By what reagents is metallic gold precipitated from solutions of its salts?

381. How is Purple of Cassius prepared?

382. Whence is platinum obtained?
383. Why are platinum utensils peculiarly adapted for use in chemical laboratories?
384. How is perchloride of platinum prepared?
385. Name the chief tests for platinum.
386. What is "platinum black"?
387. Describe an experiment demonstrative of the large amount of attraction for gases possessed by metallic platinum.
388. How is "spongy platinum" produced?
389. By what process may the metal be recovered from platinum residues?
390. What is *occlusion* in chemistry?
391. In what condition does cadmium occur in nature?
392. By what process may Iodide of Cadmium be prepared? and in what form is it used in medicine?
393. Mention the chief test for cadmium.
394. Distinguish sulphide of cadmium from other sulphides of similar color.
395. How is cadmium separated from zinc?
396. How does bismuth occur in nature?
397. What is the quantivalence of bismuth?
398. Write down equations descriptive of the actions of nitric acid on bismuth, and water on nitrate of bismuth.
399. How may pure salts be prepared from bismuth containing arsenicum?
400. Give a diagram of the process for the so-called Carbonate of Bismuth,
401. Write formulæ showing the accordance of the official Subnitrate and Carbonate with the other salts of Bismuth, and with ordinary Nitrates and Carbonates.
402. How is *Liquor Bismuthi et Ammoniæ Citratis* prepared?
403. What are the tests for Bismuth?

---

*Practical Analysis.*

Bismuth is the last of the metals whose synthetical or analytical relations are of general interest. The position of the rarer among the common metals, and the influence which either has on the other during the manipulations of analysis, will now be considered. These objects will be best accomplished, and a more intimate acquaintance with all the metals be obtained, by analyzing, or studying the methods of analyzing, solutions containing one or more metallic salts.

Of the following Tables, the first (1) includes directions for the analysis of an aqueous or only slightly acid solution, containing but one salt of any of the metals hitherto considered. Here the color of the precipitate or precipitates afforded by a metal under given circumstances must be relied on to a considerable extent in attempting the detection of the various elements.

The folded Table (2) is intended as a chart for the analysis of solutions containing salts of more than one of the common and rarer metals. It is simply a compilation from the foregoing reactions—an extension of the scheme for the analysis of salts of the ordinary metals. Hence it often may be altered or varied in arrangement to suit the requirements of the analyst.

That on p. 259 is a mere outline of the other two Tables. It gives the position of the metals in relation to each other, and will much aid the memory in recollecting that relation.

The analysis of solutions containing only one metal will, as already stated, serve to impress the memory with the characteristic tests for the various metals and other radicals, and familiarize the mind with chemical principles. Medical students seldom have time to go further than this. Much thorough analytical and general chemical knowledge is only acquired by working on such mixtures of bodies as are met with in actual practice, beginning with solutions which may contain any or all of the members of a group (see previous pages), then examining solutions containing more than one group, and finally analyzing liquids in which dissolved several salts of any of the common or rarer metals.

The Author cannot too strongly recommend students thoroughly to master the art of analysis, not only on account of its direct value, but because practice enables the learner to acquire rapidly and soundly a good knowledge of chemistry, and to improve his general mental faculties to an important extent.

**1. TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS OR ONLY SLIGHTLY ACID SOLUTION OF ORDINARY SALTS OF ONE OF THE ELEMENTS HERETO CONSIDERED—THE COMMON AND RARER METALS.**

Add hydrochloric acid.

**Precipitate** Hg(ous) Pb Ag. Collect, wash, and add AmHO.
Hg ppt., blackened.
Pb ppt., still white.
Ag ppt., dissolved.
Sb and Bi may also be precipitated by HCl, but are dissolved on adding more HCl.

**If HCl gave no precipitate the metal is still in the liquid; pass $H_2S$ through the solution.**

**Precipitate** Cd Cu Hg(ic) Pb Bi As Sb Sn Au Pt. Collect wash, add AmHS.

| Insoluble | | Soluble | |
|---|---|---|---|
| Cd, yellow. | | As(ous) | yellow. |
| Cu | black. | Sb(ic) | yellow. |
| Hg(ic) | black. | Sb, orange. | |
| Pb | black. | Sn(ous) | black. |
| Bi | black. | Au | black. |
| | | Pt | black. |

Apply special tests for each to the original solution. For these, see the previous pages.

**If $H_2S$ gave no precipitate add AmCl, AmHO, and AmHS.**

**Precipitate** Zn Mn Co Ni Al Fe Cr.
Zn, Al } white.
Cr, green.
Mn, skin-tint.
Ni, Co, Fe } black.
Test specially for each in original solution. See previous pages.

**If AmHS gave no precipitate add $Am_2CO_3$.**

**Precipitate** Ba Sr Ca. Collect, wash, dissolve in $HC_2H_3O_2$, add $K_2CrO_4$.

Ppt. Ba.

Sol. Sr. Ca. Add $H_2SO_4$.

Ppt. Sr. Sol. Ca.

**If Am... gave no precipitate add $Am_2HAsO_4$.**

Ppt. Mg.

If no precipitate test original solution in flame on loop of Pt wire.
L, crimson.
Na, yellow.
K, violet.
If neither, test orig. sol. for Am.

## 2. TABLE OF SHORT DIRECTIONS FOR APPLYING SOME OF TH

Add hydrochloric acid.

| Precipitate<br>Hg(ous) Pb Ag.<br>Wash, boil with water, filter. | Cd C |
|---|---|

| Precipitate<br>Hg(ous) Ag.<br>Wash, add Am[illegible] | Filtrate<br>Pb.<br>Add $H_2SO_4$.<br>White ppt. |
|---|---|

| Ppt.<br>Hg (ous).<br>Black ppt. | Filtrate<br>Ag.<br>Add $HNO_3$.<br>White ppt. |
|---|---|

See also p. 260.

| Precipi<br>Cd Cu Hg(ic) Pb<br>Collect, wash, digest |
|---|

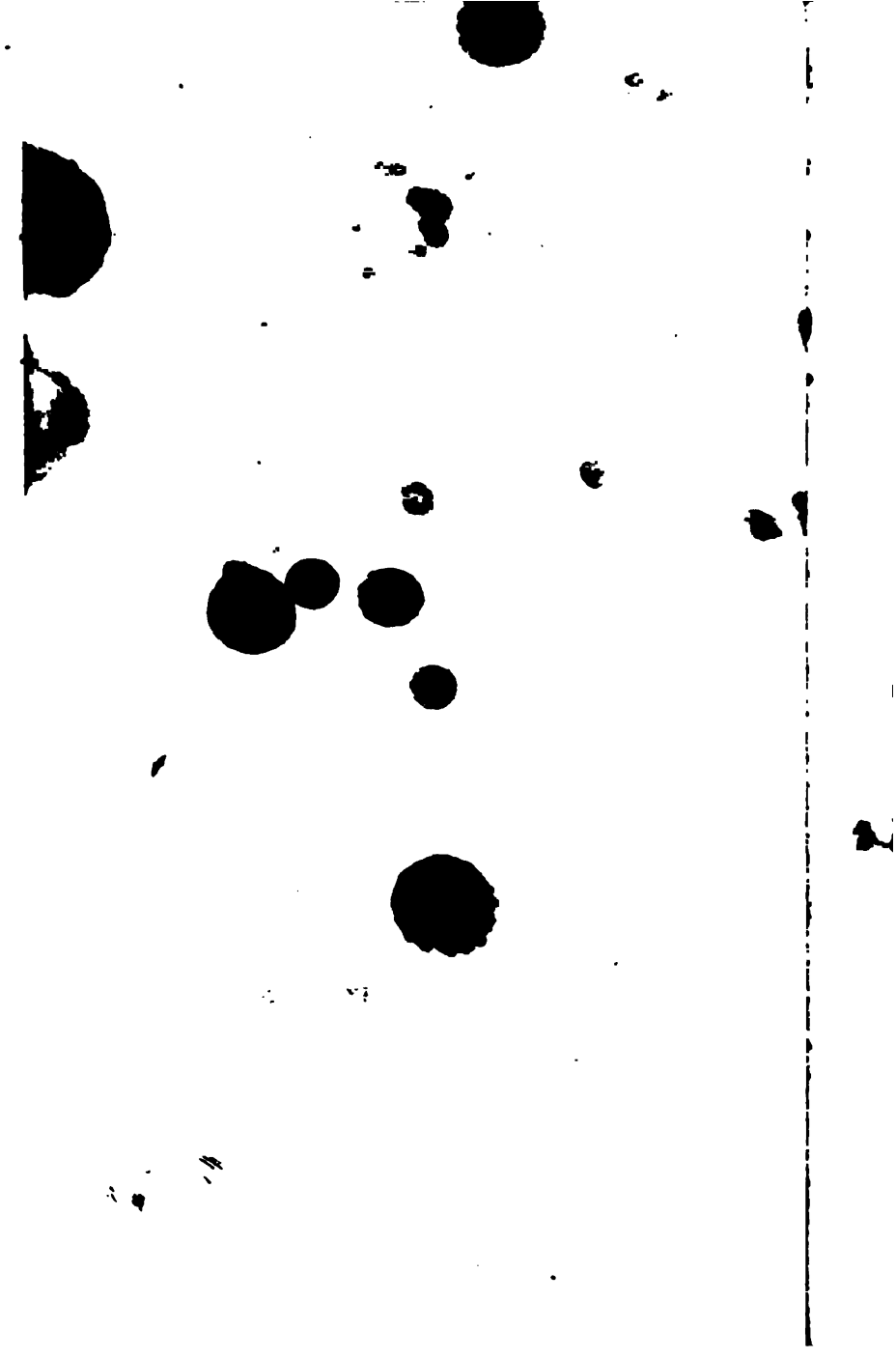
| Precipitate<br>Cd Cu Hg(ic) Pb Bi.<br>Wash, boil in $HNO_3$, filter. | Ad |
|---|---|

| Ppt.<br>Hg (ic).<br>Black.<br>Confirm by Cu test in original solution. | Filtrate<br>Cd Cu Pb Bi.<br>Add AmHO, filter. | Ins<br>As<br>Yell<br>Confi<br>by F<br>man<br>tes |
|---|---|---|

| Precipitate<br>Pb Bi.<br>Wash, add a few drops $HNO_3$, dilute, filter. | Filtrate<br>Cd Cu.<br>Add K[illegible]<br>H[illegible] |
|---|---|

| Ppt.<br>Bi. | Filt.<br>Pb.<br>Dilute, add $H_2SO_4$, set aside; white ppt. | Ppt.<br>Cd.<br>Yellow. | Filt.<br>Cu.<br>Acidify with $HC_2H_3O_2$.<br>Brown ppt. | Au a |
|---|---|---|---|---|

See also pp. 260–1.



### 3. OUTLINE OF THE ANNEXED ANALYTICAL TABLES.

| HCl | $H_2S$ | | AmHS | | $Am_2CO_3$ | $Am_2HAsO_4$ | |
|---|---|---|---|---|---|---|---|
| Hg (as mercurous salt) | Cd | Insoluble in AmHS. | Zn | Hydrates soluble in AmHO. | Ba | Mg | K |
| Pb (partially) | Cu | | Mn* | | Sr | | Na |
| | Hg (as mercuric salt) | | Co | | Ca | | Am |
| Ag | Pb (entirely) | | Ni | | | | |
| | Bi | | | | | | |
| | As (as arsenious or arsenic salt) | Soluble in AmHS. | Al | Hydrates insol. in AmHO. | | | |
| | Sb | | Fe | | | | L |
| | Sn (as stannous or stannic salt) | | Cr | | | | |
| | Au | | | | | | |
| | Pt | | | | | | |

* See page 232.

The laboratory student should practise the examination of aqueous solutions of salts of the above metals until able to analyze with facility and accuracy.

---

## MEMORANDA RELATING TO THE PRECEDING ANALYTICAL TABLES.

### *General Memoranda.*

These charts are constructed for the analysis of salts more or less soluble in water.——The student has still to learn how substances insoluble in water are to be brought into a state of solution; but, once dissolved, their analysis is effected by the same scheme as that just given. The Tables, especially the second (No. 2), may there-

fore be regarded as fairly representing the method by which metallic constituents of chemical substances are separated from each other and recognized.——The methods of isolation of the complementary constituent of the salt (the reactions of non-metals and acidulous radicals) will form the next object of practical study.

The general memoranda given in connection with the first Table (p. 224) are equally applicable to the extended second Table, and should again be carefully read through.

### *Special Memoranda.*

*The hydrochloric acid precipitate* may at first include some antimony and bismuth as oxychlorides, readily dissolved, however, by excess of acid.——If either of these elements be present, the washings of the precipitate will probably be milky; in that case add a few drops of hydrochloric acid, which will clear the liquid and make way for the application of the test for lead.

*The sulphuretted hydrogen precipitate* may be white, in which case it is nothing but sulphur; for, as already indicated, ferric salts are reduced to ferrous, and chromates to the lower salts of chromium by sulphuretted hydrogen, sulphur being deposited:—

$$Fe_2Cl_6 + 2H_2S = 4FeCl_2 + 4HCl + S_2;$$
$$4H[illegible]_4 + 6H_2S + 12HCl = 2Cr_2Cl_6 + 16H_2O + 3S_2.$$

But the precipitate may also be colored, or even white when only lead or mercury is present, through an insufficiency of sulphuretted hydrogen having produced a peculiar oxysulphide or hydrato-sulphide. The gas should be passed through the liquid until, even after well shaking, it smells strongly of sulphuretted hydrogen.

*The portion of the sulphuretted-hydrogen precipitate dissolved by sulphydrate of ammonium* may include a trace of copper, sulphide of copper being not altogether insoluble in sulphydrate of ammonium.——On adding hydrochloric acid to the sulphydrate-of-ammonium solution, a white precipitate of sulphur only may be precipitated, the sulphydrate of ammonium nearly always containing free sulphur.——Strong hydrochloric acid does not readily dissolve small quantities of sulphide of antimony out of much sulphide of arsenicum; and, on the other hand, the strong hydrochloric acid takes into solution a small quantity of sulphide of arsenicum if much sulphide of antimony is present. The precipitate or the original solution should therefore be examined by the other (hydrogen) tests for these elements if doubt exists concerning the presence or absence of either. Tin remains in the hydrogen-bottle in the metallic state, deposited as a black powder on the zinc used in the experiment. The contents of the bottle are turned out into a dish, ebullition continued until evolution of hydrogen ceases, and the zinc is taken up by the excess of sulphuric acid employed; any tin is then filtered out, washed, dissolved in a few drops of hydrochloric acid, and the liquid tested for tin by the usual reagents.——Tin may be detected in the mixed sulphides of tin, arsenicum, and antimony by the blowpipe reaction (*vide* Index).

*The portion of the sulphuretted-hydrogen precipitate not dis-*

*solved by the sulphydrate of ammonium* may leave a yellow semi-fused globule of sulphur on boiling with nitric acid. This globule may be black, not only from presence of mercuric sulphide, but also from inclosed particles of other sulphides protected by the sulphur from the action of the acid. It may also contain sulphate of lead, produced by the action of nitric acid on sulphide of lead. In cases of doubt the mass must be removed from the liquid, boiled with nitric acid till dissolved, the solution evaporated to remove excess of acid, and the residue examined; but usually it may be disregarded. ——Before testing for bismuth, any considerable excess of acid should be removed by evaporation, and the residual liquid should be freely diluted. If no precipitate (oxynitrate of bismuth) appear, chloride-of-ammonium solution may be added, oxychloride of bismuth more readily forming than even oxynitrate.——In testing for lead by sulphuric acid the liquid should be diluted and set aside for some time.

*Mercury* may also be isolated by digesting the sulphuretted-hydrogen precipitate in sulphydrate of sodium instead of sulphydrate of ammonium. The sulphides of arsenicum, antimony, tin, and mercury are thus dissolved out. The mixture is then filtered, excess of hydrochloric acid added to it, and the precipitated sulphides collected on a filter, washed, and digested in sulphydrate of ammonium; sulphide of mercury remains insoluble, while the sulphides of arsenicum, antimony, and tin are dissolved. By this method copper also appears in its right place only, sulphide of copper being quite insoluble in sulphydrate of sodium. The other metals are then separated in the usual way.

*The sulphydrate of ammonium precipitate* may, if the original solution was acid, contain Phosphates, Oxalates, Silicates, and Borates of Barium, Calcium, and Magnesium. These will subsequently come out with the iron, and, being white, give the iron precipitate a light-colored appearance; their examination must be conducted separately, by a method described subsequently in connection with the treatment of substances insoluble in water.——The precipitates containing aluminium, iron, and chromium hydrates often contain some manganese. This manganese may be detected by washing the hydrates to remove all traces of chlorides, boiling with nitric acid, adding either puce-colored oxide of lead or red lead, and setting the vessel aside. If manganese be present a red or purple liquid is produced.——Sulphide of nickel is not easily removed by filtration (*vide* p. 236) until most of the excess of sulphydrate of ammonium has been dissipated by prolonged ebullition.

*The carbonate-of-ammonium precipitate* may not contain the whole of the barium, strontium, and calcium in the mixture, unless free ammonia be present; for the carbonates of those metals are soluble in water charged with carbonic acid. If, therefore, the liquid is not distinctly ammoniacal, solution of ammonia should be added. ——Neither carbonate nor hydrate of ammonium wholly precipitates magnesian salts; and, as a partial precipitation is undesirable, a solvent, in the form of an alkaline salt (chloride of ammonium), if not already in the liquid, should be added.——In the chart opposite p. 258 strontium is ordered to be separated from calcium by adding to

the acetic solution diluted sulphuric acid. The latter, unless extremely dilute, may precipitate calcium. Any such loss of calcium is in itself of little consequence, because enough sulphate of calcium remains in the filtrate to afford a calcium reaction when ammonia and oxalate of ammonium are subsequently added. But the calcium precipitated by the sulphuric acid may be wrongly set down as strontium. Therefore test a little of the acetic solution for strontium by an aqueous solution of sulphate of calcium, when, if no precipitate falls after setting aside for several minutes, strontium may be regarded as absent. If a precipitate occurs strontium is present; the rest of the acetic solution is then tested for calcium as directed in the chart, the final testing by oxalate of ammonium being, of course, preceded by the addition of ammonia.

*Lithium.*—The search for lithium may usually be omitted. Should a precipitate, supposed to be due to lithium, be obtained, it must be tested in a flame (= scarlet tint), as a little magnesium not unfrequently shows itself under similar circumstances.

*Spectral Analysis.*—If present only in minute proportions, the lithium may also remain with the alkalies; it can then only be detected by physical analysis (by a prism) of the light emitted from a tinged flame—by, in short, an instrument termed a spectroscope. Such a method of examination is called spectral analysis, a subject of much interest and of no great difficulty, but scarcely within the range of Pharmaceutical Chemistry; it will be briefly described in connection with the methods of analyzing solid substances.

## QUESTIONS AND EXERCISES.

404. Describe a general method of analysis by which the metal of a single salt in a solution could be quickly detected.

405. Give illustrations of black, white, light pink, yellow, and orange sulphides.

406. Mention the group-tests generally employed in analysis.

407. Under what circumstances may a hydrochloric precipitate contain antimony or bismuth?

408. If a sulphuretted-hydrogen precipitate is white, what substances are indicated?

409. Give processes for the qualitative analysis of liquids containing the following substances:—

*a*. Arsenicum and Cadmium.
*b*. Bismuth and Antimony.
*c*. Ferrous and Ferric salts.
*d*. Aluminium, Iron, and Chromium.
*e*. Arsenicum, Antimony, and Tin.
*f*. Lead and Strontium.
*g*. Iron, Sodium, and Arsenicum.
*h*. Mercury, Manganese, and Magnesium.
*i*. Zinc, Manganese, Nickel, and Cobalt.
*j*. Barium, Strontium, and Calcium.

# THE ACIDULOUS RADICALS.

*Introduction.*—The twenty-six radicals which have up to this point mainly occupied attention are (admitting ammonium, $NH_4$) metals; and they have been almost exclusively studied not in the free state, but in the condition in which they exist in salts. Moreover these metals have been treated as if they formed the more important constituent, the stronger half, the foundation or base, of salts. Attention has been continuously directed to the metallic or *basylous* side of salts. And indeed there is still one more basylous radical which must be mentioned, though it is usually supposed to play only a subordinate part in medicine—Hydrogen. Unlike the salts of most metals, those of hydrogen (the so-called acids) are never, in medicine or the arts generally, professedly used for the sake of their hydrogen, but always for the other half of the salt, the *acidulous* side. And it is not for their basylous radical that these hydrogen salts are now commended to notice,* but in order to study, under the most favorable circumstances, these acidulous groupings which have continually presented themselves in operations on salts, but which were for the time of secondary importance. These *acidulous radicals* may now be treated as the primary object of attention; and there is no better way of doing so than in operating on their compounds with hydrogen, the relatively inferior medicinal importance of which element, as compared with potassium, iron, and other basylous radicals, will serve to give the desired prominence to the *acidulous radicals* in question.

*Common Acids.*—These salts of hydrogen (hydrogen easily displaceable, or in certain cases, in part, by ordinary metals) are the ordinary sharp, sour bodies termed *acids* (from the Latin root *acies*, an edge). The following Table includes the formulæ and usual names of the most important; others will be noticed subsequently. A few of those mentioned are unstable or somewhat rare; in such cases a

* It must not be forgotten that the commonest salt of any radical whatsoever is a salt of hydrogen, the oxide of hydrogen ($H_2O$), or hydrate of hydrogen (HHO), *water*. In the reactions already performed, the value of this compound has been constantly recognized, both for its hydrogen and for its oxygen, but most of all as the vehicle or medium by which nearly all other atoms are enabled to come into that contact with each other without which their existence would be almost useless; for some atoms are like some animals, out of water they are as inactive as fishes. It is true that both fishes and salts have usually to be removed from water to be utilized by man; but before they can be assimilated, either as food or as medicine, they must again seek the agency of water in becoming dissolved.

common metallic salt containing the acidulous radical may be used for reactions.

| | |
|---|---|
| $HCl$ | hydrochloric acid. |
| $HBr$ | hydrobromic acid. |
| $HI$ | hydriodic acid. |
| $HCN$ (HCy) | hydrocyanic acid. |
| $HNO_3$ | nitric acid. |
| $HClO_3$ | chloric acid. |
| $HC_2H_3O_2$ | acetic acid.* |
| $H_2S$ | hydrosulphuric acid.† |
| $H_2SO_3$ | sulphurous acid. |
| $H_2SO_4$ | sulphuric acid. |
| $H_2CO_3$? | carbonic acid. |
| $H_2C_2O_4$ | oxalic acid. |
| $H_2C_4H_4O_6$ | tartaric acid. |
| $H_3C_6H_5O_7$ | citric acid. |
| $H_3PO_4$ | phosphoric acid. |
| $H_3BO_3$ | boracic acid. |

The old names are here retained for these acids, but, in studying their chemistry and chemical relations to other salts, they are usefully spoken of by such more purely chemical names as (for hydrochloric acid) chloride of hydrogen, (for nitric acid) nitrate of hydrogen, and so on—sulphate of hydrogen, tartrate of hydrogen, phosphate of hydrogen

A prominent point of difference will at once be noticed between the basylous radicals met with up to the present time and the acidulous groupings included in the above tabular list. The former are nearly all elements, ammonium only being a compound; the latter are mostly compounds, chlorine, bromine, iodine, and sulphur being the only elements. This difference will not, however, be so apparent when the chemistry of alcohols, ethers, and such bodies has been mastered, for they are all salts of *compound* basylous radicals.

*Rarer Acids.*—The above acids contain the only acidulous groupings that commonly present themselves in analysis, or in pharmaceutical operations. There are, however, several other acids (such as hypochlorous, nitrous, hypophosphorous, valerianic, benzoic, gallic, tannic, uric, hyposulphurous, hydroferrocyanic, hydroferridcyanic,

---

* The hydrogen on the acidulous side must not be confounded with the basylous hydrogen in all these hydrogen salts or acids; the two perform entirely different functions. Hydrogen in the acidulous portion is like the hydrogen in the basylous radical ammonium, it has combined with other atoms, to form a group which plays more or less the part of an elementary radical, and to which a single symbol is not unfrequently applied (Am; Cy, $\overline{A}$, O, $\overline{T}$, $\overline{C}$, etc.). Cobalt, chromium, iron, platinum, etc., resemble hydrogen in this respect in often uniting with other atoms to form definite acidulous radicals, in which the usual basylous character of the metals has for the time disappeared. In *hydrides* (p. 122) hydrogen itself is an acidulous radical.

† Synonyms: sulphydric acid and sulphuretted hydrogen.

lactic, etc.) with which it is desirable to be more or less familiar; reactions concerning these will therefore be described. Arsenious, arsenic, stannic, manganic, and chromic acids have already been treated of in connection with the metals they contain; in practical analysis they always become sufficiently altered to come out among the metals.

*Quantivalence.*—A glance at the foregoing Table is sufficient to show the quantivalence of the acidulous radicals. The first seven are clearly univalent, then follow six bivalent, leaving three trivalent.

These all combine with equivalent amounts of basylous radicals to form various salts; hence they may be termed monobasylous, dibasylous, and tribasylous radicals. The acids themselves were formerly spoken of as monobasic, dibasic, and tribasic respectively, or monobasic and polybasic, in reference to the amount of *base* (hydrates or oxides) they could decompose; but the terms are no longer definite, and hence but little used in mineral chemistry.

*Antidotes.*—The antidotes in cases of poisoning by the strong acids will naturally be non-corrosive alkaline substances, as soap and water, magnesia, common washing "soda," or other carbonates. Vinegar, lemon-juice, and weak or non-corrosive acids would be the appropriate antidotes to caustic alkalies.

*Analysis.*—The practical study of the acidulous side of salts will occupy far less time than the basylous. Salts will then be briefly examined as a whole.

*One Word of Caution.*—It is only for convenience in the division of chemistry for systematic study that salts may be considered to contain basylous and acidulous radicals, or separate sides, so to speak; for we possess no absolute knowledge of the internal arrangement of the atoms (admitting that there are such things) in the molecule of a salt. We only know that certain groups of atoms may be transferred from compound to compound in mass (that is, without apparent decomposition); hence the assumption that these groups are radicals. A salt is probably, however, a whole, having no such sides as those mentioned.

## QUESTIONS AND EXERCISES.

410. Mention the basylous radical of acids.

411. Give illustrations of univalent, bivalent, and trivalent acidulous radicals, or monobasylous, dibasylous, and tribasylous radicals.

412. What is the difference between an elementary and a compound acidulous radical?

413. Name the grounds on which salts may be assumed to contain basylous and acidulous radicals.

## HYDROCHLORIC ACID AND OTHER CHLORIDES.

Formula of Hydrochloric Acid HCl. Molecular weight* 36.5.

The acidulous radical of hydrochloric acid and of other chlorides is the element chlorine (Cl). It occurs in nature chiefly as chloride of sodium (NaCl), either solid, under the name of *rock-salt*, mines of which are not unfrequently met with, or in solution in the water of all seas. Common table-salt is more or less pure chloride of sodium in minute crystals. Chlorine, like hydrogen, is univalent (Cl'); its atomic weight is 35.5. Its molecule is symbolized thus, $Cl_2$, chloride of chlorine.

### Reactions.

### Hydrochloric Acid.

*First Synthetical Reaction.*—To a few fragments of chloride of sodium in a test-tube or small flask add about an equal weight of sulphuric acid; colorless and invisible gaseous hydrochloric acid is evolved, a sulphate of sodium remaining. Adapt to the mouth of the vessel by a perforated cork, a piece of glass tubing bent to a right angle, heat the mixture, and convey the gas into a small bottle containing a little water; solution of hydrochloric acid results.

$$\underset{\text{Chloride of sodium.}}{NaCl} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Hydrochloric acid.}}{HCl} + \underset{\text{Acid sulphate of sodium.}}{NaHSO_4}$$

*Hydrochloric Acid.*—The product of this operation is the nearly colorless and very sour liquor commonly called hydrochloric acid. When of certain given strengths (estimated by volumetric analysis) it forms *Acidum Hydrochloricum*, B. P., and *Acidum Muriaticum*, U. S. P., and *Acidum Hydrochloricum Dilutum*, B. P. (and U. S. P., specific gr. 1.038). The former has a specific gravity of 1.16 (1.1578), and contains 31.8 per cent. of real acid, the latter specific gravity, 1.052, with 10.58 per cent. of the real acid, is made by diluting 8 fluid parts of the strong acid with water until the mixture measures 26½ fluid parts. The above process is that of the Pharmacopœias, larger vessels being employed, and the gas being freed from any trace of sulphuric acid by washing. Other chlorides yield hydrochloric acid when heated with sulphuric acid; but chloride of sodium is always used, because cheap and common.

*Common yellow hydrochloric acid* is a by-product in the manufacture of carbonate of sodium from common salt, a process in which the chloride of sodium is first converted into sulphate, hydrochloric acid being liberated. This impure acid is liable to contain iron, arsenic, fixed salts, sulphuric acid, sulphurous acid, nitrous compounds, and chlorine.

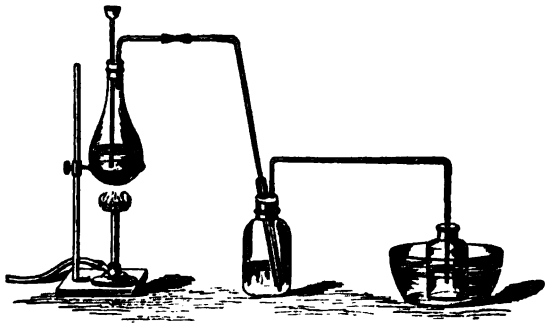
---

* The weight of a molecule is the sum of the weights of its atoms.

*The official process for the preparation of hydrochloric acid* is as follows: it may be carried out by the student with about one-twelfth of the quantities mentioned:—

"Take of chloride of sodium, dried, 48 ounces, sulphuric acid 44 fluidounces, water 36 fluidounces, distilled water 50 fluidounces; pour the sulphuric acid slowly into 32 ounces of the water, and, when the mixture has cooled, add it to the chloride of sodium previously introduced into a flask having the capacity of at least one gallon. Connect the flask by corks and a bent glass tube with a three-necked wash-bottle, furnished with a safety-tube, and containing the remaining 4 ounces of the water [or let the flask-tube pass loosely through a wider tube fixed in the cork of the wash-bottle, as shown in Fig. 37]; then, applying heat to the flask, con-

Fig. 37.



Preparation of Hydrochloric Acid.

duct the disengaged gas through the wash-bottle into a second bottle containing the distilled water, by means of a bent tube dipping about half an inch below the surface, and let the process be continued until the product measures 36 ounces, or the liquid has acquired a specific gravity of 1.16. The bottle containing the distilled water must be kept cool during the whole operation."

The modification of wash-bottle shown in the figure allows of the easy insertion or removal of a delivery-tube. The wider tube there shown, or an ordinary tube-funnel, also acts as a *safety-tube* by admitting air the moment there is any tendency in the water in the receiver to be forced back on account of a too rapid absorption of the gas. The time of the student will be saved if hydrochloric acid already in stock be placed in the wash-bottle instead of water.

*Invisible gaseous hydrochloric acid* forms visible grayish-white fumes on coming into contact with air. This is due to combination with the moisture of the air. The intense greediness of hydrochloric gas and water for each other is strikingly demonstrated on opening a test-tube full of the gas under water; the latter rushes into and instantly fills the tube. If the water is tinged with blue litmus, the

acid character of the gas is prettily shown at the same time. The test-tube, which should be perfectly dry, may be filled from the delivery-tube direct; for the gas is somewhat heavier than, and therefore readily displaces, air. The mouth may be closed by the thumb of the operator. At low temperatures hydrochloric acid and water form a crystalline compound $HCl,2H_2O$.

*Note.*—The process, as described in the British Pharmacopœia, includes the use of as much sulphuric acid as is theoretically necessary for the production of acid sulphate of sodium ($NaHSO_4$) which remains in the generating vessel. A hot solution of this residue carefully neutralized by carbonate of sodium, filtered and set aside, yields normal sulphate (*Sodii Sulphas*, U. S. P.), in the form of transparent oblique efflorescent prisms ($Na_2SO_4.10H_2O$).

$$\underset{\text{Acid sulphate of sodium.}}{2NaHSO_4} + \underset{\text{Carbonate of sodium.}}{Na_2CO_3} = \underset{\text{Sulphate of sodium.}}{2Na_2SO_4} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

## Chlorine.

*Second Synthetical Reaction.*—To some drops of hydrochloric acid (that is, the common aqueous solution of the gas) add a few grains of black oxide of manganese, and warm the mixture; *chlorine*, the acidulous radical of all chlorides, is evolved, and may be recognized by its peculiar odor, or irritating effect on the nose and air-passages.

$$4HCl + MnO_2 = Cl_2 + 2H_2O + MnCl_2$$

*Chlorine-water.*—This is the process of the Pharmacopœias for the production of chlorine-water (*Liquor Chlori*, B. P., *Aqua Chlorinii*, U. S. P.), the gas being first washed and then passed into water. One ounce of oxide to six fluidounces of acid diluted with two of water, and the gas passed through a wash-bottle containing about two ounces of water, yield enough chlorine to produce about a pint and a half of chlorine-water. (On this small scale, less than half the acid is utilized through incomplete decomposition of the materials and, especially, incomplete absorption of the chlorine gas.) Chlorine slowly decomposes water, with production of hydrochloric acid and oxygen gas; it is best preserved in a green-glass well-stoppered bottle in a cool and dark place. At common temperature (60° F.), if fresh and thoroughly saturated, chlorine-water contains more than twice (2.3) its bulk of chlorine, or less than 1 per cent. (about 0.75) by weight. Chlorine passed into cold water yields crystals of hydrous chlorine ($Cl5H_2O$), and these when heated under pressure give an upper layer of chlorine-water and a lower layer of *liquid chlorine*.

*Note.*—To obtain the chlorine from other chlorides, sulphuric acid as well as black oxide of manganese must be added. Hydrochloric acid is first formed. The action described in the above equation then goes on, except that half instead of the whole of the oxygen of the black oxide is available for the removal of the hydrogen from the chlorine of the hydrochloric acid, the other half being taken up by

the hydrogen of the sulphuric acid. Thus, supposing common salt to be the chloride used, the following equations may represent the supposed steps of the process:—

$$\begin{aligned} 2NaCl + H_2SO_4 &= Na_2SO_4 + 2HCl, \\ MnO_2 + H_2SO_4 &= MnSO_4 + H_2O \quad + O; \\ \text{then the } 2HCl + O &= H_2O + Cl_2 \end{aligned}$$

or the whole may be included in one equation:—

$$2NaCl + MnO_2 + 2H_2SO_4 = Na_2SO_4 + MnSO_4 + 2H_2O + Cl_2.$$

This reaction may have occasional analytical interest, a very small quantity of combined chlorine being recognized by its means. But the following test is nearly always applicable for the detection of this element, and leaves nothing to be desired in point of delicacy.

### *Analytical Reactions (Tests).*

To a drop of hydrochloric acid, or to a dilute solution of any other chloride, add solution of nitrate of silver; a white curdy precipitate falls. Pour off most of the supernatant liquid, add nitric acid and boil; the precipitate does not dissolve. Pour off the acid, and add dilute ammonia: the precipitate quickly dissolves. Neutralize the solution by an acid; chloride of silver is once more precipitated.

The formation of this white precipitate, its appearance, insolubility in boiling nitric acid, solubility in ammonia, and reprecipitation by an acid, form abundant evidence of the presence of chlorine. Its occurrence as a chloride of a metal is determined by testing for the metal with the appropriate reagent; its occurrence as hydrochloric acid is considered to be indicated by the odor, if strong, and the sour taste, if weak, of the liquid, and the action of the liquid on blue litmus-paper, which, like other acids, it reddens. If hydrochloric acid be present in excessive quantity it will, in addition to the above reactions, give rise to strong effervescence on the addition of a carbonate, a chloride being formed. The chlorine in insoluble chlorides, such as calamel, "white precipitate," etc., may be detected by boiling with caustic potash, filtering, acidulating the filtrate by nitric acid, and then adding the nitrate of silver.

*Antidotes.*—In cases of poisoning by strong hydrochloric acid, solution of carbonate of sodium (common washing soda) or a mixture of magnesia and water may be administered as an antidote.

---

## QUESTIONS AND EXERCISES.

414. A specimen of official Hydrochloric Acid contains 31.8 per cent. by weight of gas, and its specific gravity is 1.16; work out a sum showing what volume of it will be required, theoretically, to mix with black oxide of manganese for the production of one gallon of

chlorine-water, one fluidounce of which contains 2.66 grains of chlorine. Ans. 5½ fluidounces, nearly (5.4).

415. Why does hydrochloric acid gas give visible fumes on coming into contact with air?

416. How much chloride of sodium will be required to furnish one pound of chlorine?

417. Give the analytical reactions of chlorides.

418. What antidotes may be administered in cases of poisoning by hydrochloric acid?

---

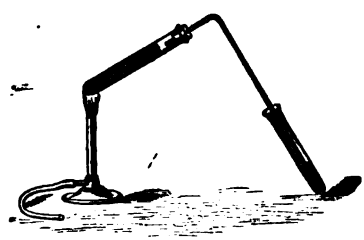
## HYDROBROMIC ACID AND OTHER BROMIDES.

Formula of Hydrobromic Acid HBr. Molecular weight 81.

*Bromine. Source, Preparation, and Properties.*—The acidulous radical of hydrobromic acid and other bromides is the element bromine, Br (*Bromum*, B. P., *Brominium*, U. S. P.). It occurs in nature chiefly as bromide of magnesium ($MgBr_2$) in sea water and certain saline springs, and is commonly prepared from the *bittern*, or residual liquors of salt-works. It may be liberated from its compounds by the process for chlorine from chlorides—that is, by heating with black oxide of manganese and sulphuric acid (see previous page). It is a dark-red volatile liquid, emitting an odor more irritating, if possible, than chlorine—of specific gravity 2.966, boiling-point 145.4° F.

*Quantivalence.*—The atom of bromine, like that of chlorine, is univalent (Br'). The atomic weight of bromine is 80. Free bromine has the molecular formula $Br_2$, bromide of bromine.

Fig. 38.



Preparation of Hydrobromic Acid.

*Hydrobromic Acid.*—The bromide of hydrogen, hydrobromic acid, may be made by decomposing bromide of phosphorus by water—$PBr_5 + 4H_2O = 5HBr + H_3PO_4$. A small quantity may be prepared by placing seven or eight drops of bromine at the bottom of a test-tube, putting in fragments of glass to the height of about an inch or two, then ten or eleven grains of phosphorus, then another inch of glass, and finally a couple of inches of glass fragments slightly wetted

with water, a delivery-tube being fitted on by a cork. The phosphorus combines readily, almost violently, with the bromine as soon as the vapor of the latter, aided by a little warmth from a flame, rises to the region of the phosphorus. The bromide of phosphorus thus formed then suffers decomposition by the water of the moist glass, phosphoric and phosphorous acids being produced. The hydrobromic acid gas passes over (heat being applied in the after part of the operation), and may be condensed in water or in solution of ammonia. The latter solution on evaporation yields bromide of ammonium.

*Other Methods.*—One hundred parts of sodium hyposulphite, fifty of bromine, and ten of water are placed in a flask and the generated gas is conducted into the upper portion of 140 parts of water contained in another vessel. When the gas begins to come over slowly, gentle heat is applied. The product is nearly 190 parts of liquid containing 25 per cent. of real acid; specific gravity 1.204. It should be kept in a cool dark place (Hager). Squibb prefers to decompose solution of bromide of potassium by sulphuric acid, and, after removal of potassium sulphate by crystallization, to distil the residual fluid. Wade prescribes an almost pure clear solution of the acid made by shaking together 120 grains of bromide of potassium and 153 grains of crystallized tartaric acid in 1 ounce of distilled water, and setting aside till precipitation of acid tartrate of potassium ceases.

*Bromide of Potassium* (KBr) is occasionally employed in pharmacy, and is the salt, therefore, which may be used in studying the reactions of this acidulous radical. The official method of making the salt has been alluded to under the salts of potassium (page 76).

*Other Bromides* are seldom used; they may be prepared in the same way as, and closely resemble, the corresponding chlorides or iodides. *Bromide of Sodium* crystallizes in anhydrous cubes (NaBr) from solutions at 110° or 120° F., and in hydrous prisms ($NaBr,2H_2O$) at ordinary temperatures.

*Bromide of Ammonium* (AmBr) (*Ammonii Bromidum*, B. P. and U. S. P.) is prepared by agitating iron wire with a solution of bromine until the odor of bromine can be no longer perceived, adding solution of ammonia, filtering, and evaporating the filtrate to dryness. It forms a white granular salt, which becomes slightly yellow on exposure to air, is readily soluble in water, less so in spirit, and, when heated, sublimes. Bromide or Iodide of Ammonium may also be made by mixing equivalent quantities of strong hot aqueous solutions of the corresponding potassium salts and of sulphate of ammonium. To the cooled liquids rectified spirit is added, which precipitates the sulphate of potassium. The spirit recovered by distillation of the clear liquid leaves the required salt as a residue in the retort.

*Solution of Bromine*, B. P., 10 minims in 5 ounces, is an aqueous solution, bromine being slightly soluble in water.

*Hypobromites*, *Bromates*, *Perbromates*, analogous to hypochlorites, chlorates, and perchlorates, are producible.

*Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a few drops of solution of a bromide (KBr, or $NH_4Br$) add solution of nitrate of silver; a yellowish-white precipitate of bromide of silver (AgBr) falls. Treat the precipitate successively with nitric acid and dilute ammonia, as described for the chloride of silver; it is only sparingly dissolved by the ammonia.

*Second Analytical Reaction.*—To solution of a bromide add a drop or two of chlorine-water, or a bubble or two of chlorine gas; then add a few drops of chloroform or ether, shake the mixture, and set the test-tube aside; the chlorine, from the greater strength of its affinities, liberates the bromine, which is dissolved by the chloroform or ether, the solution falling to the bottom of the tube in the case of the heavy chloroform, or rising to the top in the case of the light ether. Either solution has a distinct yellow or reddish-yellow or red color, according to the amount of bromine present.

*Note.*—This reaction serves for the isolation of bromine when mixed with many other substances. Excess of chlorine must be avoided, as colorless chloride of bromine is then formed. Iodides give a somewhat similar result; the absence of iodine must therefore be insured by a process given in the next section. The above solution in chloroform or ether may be removed from the tube by drawing up into a *pipette* (small pipe, a narrow glass tube, usually having a bulb or expanded portion in the centre), the bromide fixed by the addition of a drop of solution of potash or soda, the chloroform or ether evaporated off, and the residue tested as described in the next reaction.

The above operation is frequently employed for synthetical purposes.

*Third Analytical Reaction.*—Liberate bromine from a bromide by the cautious addition of chlorine or chlorine-water, then add a few drops of cold decoction of starch; a yellow combination of bromine and starch, commonly termed "bromide of starch," is formed.

*Decoction of starch* is made by rubbing down two or three grains of starch with some drops of cold water, then adding much more water and boiling the mixture.

*The above reaction may be varied* by liberating the bromine by a little black oxide of manganese and a drop of sulphuric acid, the upper part of the inside of the test-tube being smeared over with some thick decoction of starch or thin starch-paste. Even sulphuric acid alone, if strong, liberates bromine from a bromide, the hydrogen of the hydrobromic acid

first produced uniting with oxygen of the sulphuric acid—the latter being reduced to sulphurous acid or even to hydrosulphuric acid.

## HYDRIODIC ACID AND OTHER IODIDES.

Formula of Hydriodic Acid HI. Molecular weight 128.

*Source.*—The acidulous radical of hydriodic acid and other iodides is the element iodine (I). It occurs in nature chiefly as iodide of sodium and of magnesium in sea-water. Seaweeds, sponges, and other marine organisms, which derive much of their nourishment from sea-water, store up iodides in their tissues, and it is from the ashes of these that supplies of iodine (*Iodum*, B. P., *Iodinium*, U. S. P.) are obtained.

*Process.*—The seaweed ash or *kelp* is treated with water, insoluble matter thrown away, and the decanted liquid evaporated and set aside to allow of the deposition of most of the sulphates, carbonates, and chlorides of sodium and potassium. The residual liquor is treated with excess of sulphuric acid, which causes evolution of carbonic and sulphurous or sulphuretted gases, deposition of sulphur and more sulphate of sodium, and formation of hydriodic acid. To the decanted liquid is added black oxide of manganese, and the mixture is then slowly distilled; the iodine sublimes, and is afterwards purified by re-sublimation.

$$2HI + MnO_2 + H_2SO_4 = MnSO_4 + 2H_2O + I_2.$$

*The analogy of chlorine, bromine, and iodine* is well indicated by the fact that each is obtained from its compounds by the same reaction. Iodine is liberated from any iodide as bromine from bromides, or chlorine from chlorides—namely, by the action of black oxide of manganese and sulphuric acid.

*Properties.*—Iodine is a crystalline purplish-black substance; its vapor, readily seen on heating a fragment in a test-tube, is dark violet. Its vapors are irritating to the lungs; but a trace may be inhaled with safety (*Vapor Iodi*, B. P.). It melts at 239°, boils at about 392° F., and is entirely volatilized, the first portions containing any cyanide of iodine that may be present. The latter body occurs in slender colorless prisms, emitting a pungent odor.

*Quantivalence.*—The atom of iodine, like those of bromine and chlorine, is univalent* (I′). The atomic weight of iodine is 127, its molecular formula $I_2$.

---

* There is a compound of iodine having the formula $ICl_3$. Iodine would at first sight, therefore, seem to be a trivalent element (I‴); and bromine and fluorine, from their close chemical analogy with iodine, would necessarily be regarded as trivalent also. From this aspect the position of chlorine would be anomalous. Possibly, however, the compound is only a molecular combination of chloride of iodine, ICl, with chlorine, $Cl_2$. Iodine also forms with iodide of potassium a periodide, or tri-iodide, $KI_3$, which may be obtained in lustrous

*The Iodide of Hydrogen, or Hydriodic Acid*, is a heavy, colorless gas. Its solution in water may be made by passing sulphuretted hydrogen through water in which iodine is suspended.

$$2H_2S + 2I_2 = S_2 + 4HI.$$

Kolbe prepares hydriodic acid by adding to 10 parts of iodine contained in a flask, in an atmosphere of carbonic acid gas, 1 part of phosphorus, little by little and slowly. On the mixture of di- and tri-iodide of phosphorus are poured 4 parts of water. From this abundance of hydriodic acid is evolved on the application of gentle heat, and it is not contaminated by free iodine. Phosphoric acid remains.

*Iodide of potassium* (KI) is largely used in medicine, and hence is the most convenient iodide on which to experiment in studying the reactions of this acidulous radical. Solid iodine itself might be taken for the purpose; but its use and action in that state have already been alluded to in describing the iodides of potassium, cadmium, and mercury; its analytical reactions in the combined condition are those which may now occupy attention.

*Solution of Iodine.*—Iodine is slightly soluble in water (iodine-water), and readily soluble in an aqueous solution of iodide of potassium. Twenty grains of iodine and 30 of iodide of potassium dissolved in 1 ounce of distilled water, form *Liquor Iodi*, B. P.; *Liquor Iodinii Compositus*, U. S. P., is of the same character and about the same strength: 32 grains of iodine and 32 of iodide of potassium, rubbed with 1 fluidrachm of proof spirit, and 2 ounces of lard gradually mixed in, form *Unguentum Iodi*, B. P. *Unguentum Iodinii Compositum*, U. S. P., and *Unguentum Iodinii*, U. S. P., are similar preparations. It is more soluble in spirit (*Tinctura Iodinii*, U. S. P.), or in a spirituous solution of iodide of potassium (*Tinctura Iodi*, B. P., *Tinctura Iodinii Composita*, U. S. P.). It combines with sulphur, forming an unstable grayish-black solid iodide ($S_2I_2$), having a radiated crystalline structure (*Sulphuris Iodidum*, B. P. and U. S. P.; and *Unguentum Sulphuris Iodidi*). If 100 grains be thoroughly boiled with water, the iodine will pass off in vapor, and about 20 grains of sulphur remain.—*B. P.* and *U. S. P.*

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a few drops of an aqueous solution of an iodide (*e. g.* KI) add solution of nitrate of silver; a light yellow precipitate of iodide of silver (AgI) falls. Pour away the supernatant liquid and treat the precipitate with nitric acid, it is not dissolved; pour away the acid and then add dilute ammonia, it is only sparingly dissolved.

This reaction is useful in separating iodine from most other acidulous radicals, but does not distinguish iodine from bromine.

---

prismatic crystals. This, too, may have the formula $KI,I_2$. A mercuric hexiodide ($HgI_6$, perhaps $HgI_2,I_2,I_2$) is also known; and a periodide of ammonium, $NH_4I_2,I_2$.

Ammonia, it will be remembered, dissolves chloride of silver readily; hence the presence of chloride of potassium in bromide or iodide may be detected by dissolving in water, adding excess of nitrate of silver, collecting the precipitate, washing, digesting in ammonia, filtering, and adding excess of nitric acid to the filtrate; a white curdy precipitate indicates a chloride (of potassium). Bromide and iodide of silver are, however, slightly soluble in ammonia. A better process is given on page 276.

*Second Analytical Reaction.*—Liberate iodine from an iodide by the cautious addition of chlorine, then add cold decoction of starch; a deep-blue combination of iodine and starch, commonly termed "iodide of starch," is formed.

Starch is highly sensitive to the action of iodine; this reaction is consequently very delicate and characteristic. The reaction is not observed in hot liquids. Excess of chlorine must be avoided, or colorless chloride of iodine will be produced. Nitrous acid, or a nitrite acidulated with sulphuric acid, may be used instead of chlorine. Concentrated sulphuric acid also liberates iodine from iodides, the hydrogen of the hydriodic acid first produced uniting with the oxygen of the sulphuric acid—the latter ($H_2SO_4$) being reduced to sulphurous acid ($H_2SO_3$) or even to hydrosulphuric acid ($H_2S$).

In testing bromine for iodine the bromine must be nearly all removed by solution of sulphurous acid before the decoction of starch is added.

*Ozone* ($O_3$).—Papers soaked in mucilage of starch containing iodide of potassium form a test for free chlorine and nitrous acid, and are also employed by meteorologists to detect an allotropic or physically polymeric and energetic form of oxygen termed by Schönbein *ozone* (from ὀζω, *ozō*, I smell). This substance liberates iodine from iodide of potassium (with formation of iodide of starch), and is supposed to occur normally in the atmosphere, the salubrity or insalubrity of which is said to be dependent to some extent on the presence or absence of ozone. The possible occurrence of nitrous or chlorinoid gases in the air, however, renders the test untrustworthy. Houzeau proposes to test for ozone by exposing litmus paper of a neutral tint soaked in a dilute solution of iodide of potassium; the potash set free by action of the ozone turns the paper blue. The same paper without iodide would indicate the extent to which the effect might be due to ammonia vapor. Ozone, or rather ozonized air, is produced artificially in large quantities on passing air through a box (Beane's Ozone-generator) highly charged with electricity. In the latter operation condensation of the volume of air, or, rather, of the oxygen in the air, occurs. Small quantities are said to be obtained by exposing in a loosely closed bottle a stick of phosphorus partially covered by water, but the product is probably peroxide of hydrogen. It is a powerful bleaching, disinfecting, and general oxidizing agent; insoluble in water, soluble in oils of turpentine, cinnamon, and some other liquids. From experiments that have been made by Soret on the specific gravity of ozone, its molecular formula would seem to be

$O_3$, that of ordinary oxygen being $O_2$. Its smell is peculiar. (See p. 238, also "Blood.")

*Third Analytical Reaction.*—To a neutral aqueous solution of an iodide, add a solution containing one part of sulphate of copper to two and a half parts of green sulphate of iron, and well shake; a dirty-white precipitate of cuprous iodide ($Cu_2I_2$) falls.

$$2KI + 2CuSO_4 + 2FeSO_4 = Cu_2I_2 + K_2SO_4 + Fe_23SO_4.$$

Or to the liquid containing an iodide add the solution of copper sulphate and some solution of sulphurous acid, and warm the mixture, cuprous iodide falls.

$$2KI + 2CuSO_4 + H_2SO_3 + H_2O = Cu_2I_2 + 2KHSO_4 + H_2SO_4.$$

*Separation of Chlorides, Bromides, and Iodides.*—Chlorides and bromides are not affected in this way; the reaction is useful, therefore, in removing iodine from a solution in which chlorides and bromides have to be sought. The total removal of iodine by the former of the two modifications of the process is insured by supplementing the addition of the cupric and ferrous sulphates by a few drops of solution of potash or soda, any acid which might be keeping cuprous iodide in solution being thereby neutralized, ferric or ferrous hydrate, precipitated at the same time, not affecting the reaction. Occasionally, too, it may be necessary to repeat the process with the filtrate before the last traces of iodine are removed. The second modification of the process is, on the whole, to be preferred.

Chloride of the rare metal palladium performs a similar useful office in removing iodine, but not bromine or chlorine, from solutions.

Chlorides may be separated from bromides by taking advantage of the ready solubility of chloride of silver, and slow and slight solubility of bromide of silver in ammonia, especially in (a fair, not a great, excess of) ammonia containing chloride of silver.

Siebold's test for the detection of chlorides when occurring with bromides is based upon the fact that a strong solution of permanganate of potassium liberates the bromine from moderately strong solutions of bromides containing a large excess of sulphuric acid. A strong solution of permanganate is added to the aqueous solution of bromide or iodide (containing not more than 1 in 40), strongly acidified by dilute sulphuric acid, until the permanganate ceases to be decolorized, and a copious precipitate of oxide of manganese is formed. The mixture is boiled for about five minutes to expel the bromine, and then filtered. The colorless filtrate is now quite free from bromine, and may be tested for chlorine by nitrate of silver. If a chloride be present, a small quantity of its chlorine is lost by this process; but the main portion always remains, provided that no undue excess of the permanganate be used. It is essential, therefore, that the filtrate should be colorless; for if it be colored so as to indi-

cate the presence of undecomposed permanganate, the loss of the greater part if not of the whole of the chlorine must be expected. If, on the other hand, too little permanganate be employed, a trace of bromide may be left in the filtrate. If the solution under examination should be very much stronger than 1 in 40, water should be added before boiling (just after the addition of the permanganate), in order to avoid a loss of HCl.

Chlorides may also be detected in bromides and iodides by taking advantage of the formation of chlorochromic anhydride (page 239) and the non-occurrence of corresponding compounds of bromine or iodine.

To a solution of a chloride with a bromide and an iodide add a concentrated solution of sulphate of sodium, and then a reagent prepared by mixing equal volumes of sulphuric acid and saturated solution of sulphate of copper, until no further precipitation of cuprous iodide occurs. Next add solution of soda to remove excess of sulphate of copper; filter and evaporate to dryness. Place the dried residue, together with an equal bulk of red chromate of potassium, in a dry test-tube fitted with a delivery tube or into a small retort, and cover the mixture with sulphuric acid. Distil into water. Chromic anhydride and hydrochloric and hydrobromic acids are liberated by the sulphuric acid, and reacting upon one another form chlorochromic anhydride together with free bromine and chlorine.

$$CrO_3 + 2HCl = CrCl_2O_2 + H_2O.$$
$$2CrO_3 + 6HCl + 3H_2SO_4 = Cr_23SO_4 + 3Cl_2 + 6H_2O.$$
$$2CrO_3 + 6HBr + 3H_2SO_4 = Cr_23SO_4 + 3Br_2 + 6H_2O.$$

The chlorochromic anhydride is decomposed by the excess of water into which it falls, giving rise to chromic acid, which imparts its color to the liquid, and hydrochloric acid.

$$CrCl_2O_2 + 2H_2O = H_2CrO_4 + 2HCl.$$

The chlorine escapes and the bromine is dissolved by the water. The colored liquid is then shaken with chloroform, which removes the bromine—indicating bromine in the original substance. A yellow color remaining is due to chromic acid—indicating chlorides in the original substance. Or add ammonia to the distillate—the color due to bromine is thereby entirely removed while that of the chromic compound is only slightly modified.

*Fourth Analytical Reaction.*—Iodides have been shown to be useful in testing for mercuric salts (see the Mercury reactions, p. 196); a mercuric salt (corrosive sublimate, for example) may therefore be used in testing for iodides, a scarlet precipitate of mercuric iodide ($HgI_2$) being produced.

This reaction may be employed where large quantities of an iodide are present; but its usefulness in analysis is much impaired by the fact that the precipitate is soluble in excess of the dissolved iodide, or in excess of the mercuric reagent. Its color and insolubility in

water distinguish it from mercuric chloride, bromide, and cyanide, which are white soluble salts.

*Fifth Analytical Reaction.*—Iodides have also (see the Lead reactions, p. 212) been shown to be useful in testing for lead salts; similarly a lead salt (acetate, for example) may be used in testing for iodides, a yellow precipitate of iodide of lead ($PbI_2$), soluble in hot water and crystallizing in yellow scales on cooling, being produced.

Chloride, bromide, and cyanide of lead are white; hence the above reaction may occasionally be useful in distinguishing iodine from the allied radicals. But iodide of lead is slightly soluble in cold water; hence small quantities of iodide cannot be detected by this reaction. (For *Iodates* see p. 296.)

*Analogies between Chlorine, Bromine, Iodine, and their Compounds.*—These elements form a natural group or family, each distinct from the other yet closely related. Moreover, their dissimilarities are so curiously gradational as to irresistibly suggest the idea that some day we may find the differences between these bodies to be in degree rather than in kind. Thus chlorine is a gas and iodine a solid, while bromine occupies the intermediate condition. The atomic weight of bromine is nearly midway between those of chlorine and iodine. The same may be said of the weight of equal volumes of each in the gaseous state. The specific gravity of fluid chlorine is 1.33, of iodine 4.95, while bromine is nearly 3. Liquid chlorine is transparent, iodine opaque, bromine intermediate. The crystalline forms of the chloride, bromide, and iodide of a metal are commonly identical. One volume of either element in the gaseous state combines with an equal volume of hydrogen (at the same temperature) to form two volumes of a gaseous acid, very soluble in water (hydrochloric acid, hydrobromic acid, hydriodic acid). Many other analogies are traceable.

---

## QUESTIONS AND EXERCISES.

419. State the method by which Bromine is obtained from its natural compounds.

420. Mention the properties of bromine.

421. How may the Bromides of Potassium and Ammonium be made?

422. By what reagents may bromides be distinguished from chlorides?

423. Whence is iodine obtained?

424. By what process is iodine isolated?

425. State the properties of iodine.

426. What is the nature of Iodide of Sulphur?

427. Give the analytical reaction of iodides.

428. Which three substances may indirectly be detected by a mixture of iodide of potassium and mucilage of starch?

429. Describe a method by which iodides may be removed from a solution containing chlorides and bromides.

---

## HYDROCYANIC ACID AND OTHER CYANIDES.

Formula of Hydrocyanic Acid HCN or HCy.

Molecular weight 27.

*History of Cyanogen.*—The acidulous radical of hydrocyanic acid and other cyanides is a compound body, cyanogen (Cy). It is so named from *κύανος, kuanos*, blue, and *γεννάω, gennaō*, I generate, in allusion to its prominent chemical character of forming, with iron, the different varieties of Prussian blue. It was from Prussian blue that Scheele, in 1782, first obtained what we now, from our knowledge of its composition, term hydrocyanic acid, but which he called Prussic acid. Cyanogen was isolated by Gay-Lussac in 1814, and was the first compound radical distinctly proved to exist.

*Sources.*—Cyanogen does not occur in nature, and is only formed from its elements under certain circumstances. It is found in small quantities among the gases of iron-furnaces, and is produced to a slight extent in distilling coals for gas. In the form of ferrocyanide of potassium it is obtained abundantly by heating animal refuse containing nitrogen, such as the scrapings of horns, hoofs, and hides (5 parts) with carbonate of potassium (2 parts) and waste iron (filings, etc.) in a covered iron pot. The residual mass is boiled with water, the mixture filtered, and the filtrate evaporated and set aside for crystals to form. The cyanogen, produced from the carbon and nitrogen of the animal matter, unites with the potassium, and afterwards, on boiling with water, with iron, to form what is known as the yellow prussiate of potash (*Potassæ Prussias Flava*, B. P.), or ferrocyanide of potassium ($K'_4Fe''Cy'_6$) (*Potassii Ferrocyanidum*, U. S. P.), a compound occurring in four-sided tabular yellow crystals. It contains the elements of cyanogen, yet it is not a cyanide, for it is not poisonous, and is otherwise different from cyanides; it will be further noticed subsequently. From this salt all cyanides are directly or indirectly prepared.

*Cyanide of potassium* (KCy) (*Potassii Cyanidum*, U. S. P.), which is the most common, is procured by fusing 8 parts of ferrocyanide with 3 of carbonate of potassium in a crucible; carbonic acid gas ($CO_2$) is evolved, iron (Fe) is set free, and cyanate of potassium (KCyO), a body that will be subsequently noticed, is formed at the same time:—

$$2K_4FeCy_6 + 2K_2CO_3 = 10KCy + 2KCyO + Fe_2 + 2CO_2.$$

*Double cyanides* exist, such as the cyanide of sodium and silver (NaCy, AgCy), formed in the process (subsequently described) of

quantitatively determining the amount of hydrocyanic acid in a liquid by a standard solution of nitrate of silver; these compounds have, more or less, the properties of their constituents. But other cyanogen compounds, not double cyanides, occur in which the cyanogen is so intimately united with a metal as to form a distinct radical; such are ferrocyanides and ferridcyanides—salts which will be noticed in due course.

*Cyanogen*, like chlorine, bromine, and iodine, is univalent (Cy′). It may be isolated by simply heating mercuric cyanide ($HgCy_2$) or cyanide of silver (AgCy). It is a colorless gas, burning, when ignited, with a beautiful peach-blossom-colored flame.

*Mercuric cyanide* is produced in crystals on dissolving 1 part of ferrocyanide of potassium in 15 parts of boiling water, adding 2 parts of mercuric sulphate, keeping the whole hot for ten or fifteen minutes and then filtering and setting aside to cool. In addition to mercuric cyanide ($HgCy_2$), mercury (Hg), ferric sulphate ($Fe_2 3SO_4$), and sulphate of potassium ($K_2SO_4$), are formed. Any excess of ferrocyanide also gives Prussian blue by reaction with the ferric sulphate. It (*Hydrargyri Cyanidum*, U. S. P.) may also be made by dissolving red oxide of mercury in diluted hydrocyanic acid. A small flame of cyanogen may be obtained on heating a few crystals of mercuric cyanide in a short piece of glass tubing closed at one end, and applying a light to the other end as soon as evolution of gas commences; brown *paracyanogen* ($C_3N_3$) and mercury remain.

### Reactions.

### Diluted Hydrocyanic Acid.

*Synthetical Reaction.*—Dissolve 2 or 3 grains of ferrocyanide of potassium in 5 or 6 times its weight of water in a test-tube, add a few drops of sulphuric acid and boil the mixture, conveying the evolved gas by a bent glass tube (adapted to the test-tube by a cork) into another test-tube containing a little water; the product is a dilute solution of hydrocyanic acid. Made by this process in large quantities of a certain definite strength (2 per cent.), this solution is the *Acidum Hydrocyanicum Dilutum*, B. P. and U. S. P. "A colorless liquid of a peculiar odor. Specific gravity 0.997."

$$2K_4FeCy_6 + 6H_2SO_4 = FeK_2Fe''Cy_6 + 6KHSO_4 + 6HCy.$$

The details of the official process are as follows: Dissolve 2½ ounces of ferrocyanide of potassium in 10 ounces of water, add 1 fluidounce of sulphuric acid previously diluted with four ounces of water and cooled. Put the solution into a flask or other suitable apparatus of glass or earthenware, to which are attached a condenser and a receiver arranged for distillation (see p. 126), and having put 8 ounces of distilled water into the receiver, and provided efficient means for keeping the condenser and receiver cold, apply heat to the flask, until, by slow distillation, the liquid in the receiver

is increased to 17 fluidounces.* Add to this 3 ounces of distilled water, or as much as may be sufficient to bring the acid to the required strength, so that 100 grains (or 110 minims) of it, precipitated with a solution of nitrate of silver (*vide* paragraphs on quantitative analysis) shall yield 10 grains of dry cyanide of silver. The end of the condenser should pass quite into the receiver.

*The residue of this reaction* is acid sulphate of potassium ($KHSO_4$), which remains in solution, and ferrocyanide of potassium and iron ($Fe''K_2FeCy_6$), an insoluble powder sometimes termed Everitt's yellow salt, from the name of the chemist who first made out the nature of the reaction. The latter compound becomes bluish-green during the reaction, owing to absorption of oxygen. *Diluted hydrocyanic acid* may also be prepared by reaction of cyanide of silver and diluted hydrochloric acid.

*Pure anhydrous hydrocyanic acid* is a colorless, highly volatile, intensely poisonous liquid, solidifying when cooled to a low temperature. It may be made by passing sulphuretted hydrogen over mercuric cyanide. The official solution of the acid is fairly stable, but is said to be rendered more so by the presence of a minute trace of

* This operation is peculiarly liable to those sudden and tumultuous evolutions of vapor, or "bumpings," or "*soubresauts*," which often interfere with successful distillation. Such phenomena occur, according to Tomlinson, whenever unaided heat has to overcome the great amount of adhesion naturally existing between certain liquids and vapors, or, rather, between the normal liquid and those particles of it which, becoming strongly heated at the heated part of the vessel, have assumed the condition of particles of dissolved vapor, and which would at once pass from this condition into that of permanent vapor but for adhesion. Ordinarily a glass or other surface is not absolutely clean, but is more or less covered with specks, traces of materials deposited from the air, the fingers, cloths, etc. Some liquids seem to have little or no adhesion for these materials, while certain vapors have greater adhesion for the films than for the liquids. Hence, in ordinary regular ebullitions the vapors accumulate on the films, and then at once become subject to the pressure of the mass of fluid, and so pass off in bubbles. But when the films are absent, or have become removed during distillation, the heat accumulates until it is sufficient to overcome the adhesion of the super-heated particles, and these are then, all of them at once, converted into vapor, the liquid commonly boiling over, sometimes even bursting the vessel. "Bumping" would be prevented by the introduction of fragments of substances for which vapor-particles have adhesion, but no known substance has this property in an absolute degree. Fragments of tobacco-pipe or pumice-stone, pieces of cork, thick paper, resin, sulphur, platinum wire, etc., are all useful when there is no chemical action between them and the liquid. Mr. Tomlinson very strongly recommends cocoanut shell charcoal to be used whenever practicable. A slow current of gas, such as hydrogen, air, or carbonic acid gas, also usefully promotes escape of vapor from a liquid. When the bumping cannot well be prevented, as in the distillation of sulphuric acid, it is somewhat reduced in violence if the retort be heated by an annular gas-burner rather than by a single central jet.

sulphuric or hydrochloric acid. A stronger acid is liable to assimilate the elements of water, and yield formiate of ammonium ($NH_4CHO_2$). Solutions of hydrocyanic acid often become brown by formation of what is, apparently, *paracyanogen* ($C_3N_3$). According to Williams aqueous hydrocyanic acid containing 20 per cent. of glycerine can be kept for an apparently indefinite length of time.

*Note.*—A few drops of diluted hydrocyanic acid so placed that its vapor may be inhaled, forms the *Vapor Acid Hydrocyanici*, B. P., or Inhalation of Hydrocyanic Acid.

Hydrocyanic acid also occurs in cherry-laurel water and bitter almond water (*vide* Index).

The methods of determining the strength of solutions of hydrocyanic acid will be described in connection with volumetric and gravimetric quantitative analysis. They are based on the formation of cyanide of silver and its solubility in solution of cyanide of potassium, as described in the next reaction.

The hydrocyanic acid used in pharmacy is extremely liable to variation in strength. It should frequently be tested volumetrically.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a few drops of the hydrocyanic acid solution produced in the above reaction, or to any solution of a cyanide, add excess of solution of nitrate of silver; a white precipitate of cyanide of silver (AgCy) falls. When the precipitate has subsided, pour away the supernatant liquid, and place half of the residue in another test-tube: to one portion add nitric acid, and notice that the precipitate does not dissolve; to the other add ammonia, and observe that the precipitate is insoluble or only sparingly soluble. (Chloride of silver, which is also white, is readily soluble in ammonia.) Cyanide of silver dissolves in solutions of cyanides of alkali-metals, soluble double cyanides being formed (*e. g.*, KCy, AgCy).

*Solubility of precipitates in strong solutions of salts.*—Cyanide of silver and many other pricipitates insoluble in acids (similar remarks apply to precipitates insoluble in alkalies) are often soluble in the strong saline liquids formed by the addition of acids and alkalies to one another. Hence the precaution of adding the latter reagents to separate portions of a precipitate, or of not adding the one until the other has been poured away.

*Cyanogen in an insoluble cyanide*, such as cyanide of silver itself, is readily recognized on heating the substance in a short piece of glass tubing closed at one end like a test-tube and drawn out at the other end, so as to have but a small opening; on applying a flame, the escaping cyanogen ignites and burns with a characteristic peach-blossom tint.

## Antidote.

*Second Analytical Reaction.*—To a dilute solution of hydrocyanic acid, or a soluble cyanide, add a few drops of solution of a ferrous salt and a drop or two of solution of a ferric salt (ferrous sulphate and ferric chloride are usually at hand); to the mixture add potash or soda (magnesia or carbonate of sodium), and then hydrochloric acid; a precipitate of Prussian blue remains. The decompositions may be traced in the following equations:—

$$\begin{array}{llll} HCy & + KHO & = KCy & + H_2O \\ 2KCy & + FeSO_4 & = FeCy_2 & + K_2SO_4 \\ 4KCy & + FeCy_2 & = K_4FeCy_6 & \text{or } K_4Fcy \\ 3K_4Fcy & + 2Fe_2Cl_6 & = 12KCl & + Fe_4Fcy_3. \end{array}$$

The test depends on the conversion of the cyanogen into ferrocyanogen by aid of the iron of a ferrous salt, and the combination of the ferrocyanogen, so produced, with the iron of a ferric salt.

Hence a mixture of green sulphate of iron, solution of perchloride of iron, and either magnesia or carbonate of sodium, is the recognized *antidote* in cases of poisoning by hydrocyanic acid or cyanide of potassium.

In such an alkaline mixture the poisonous cyanide, by reaction with ferrous hydrate, is at once converted into innocuous ferrocyanide of potassium or sodium, etc.; should the mixture become acid, the ferric salt present reacts with the soluble ferrocyanide, forming insoluble Prussian blue which is also inert. From the rapidity of the action of these poisons, however, there is seldom time to prepare an antidote. Emetics, the stomach pump, the application of a stream of cold water to the spine, and the above antidote, form the usual treatment.

*Third Analytical Reaction.*—To solution of hydrocyanic acid add ammonia, and common yellow sulphydrate of ammonium, and evaporate the liquid nearly or quite to dryness in a small dish, occasionally adding ammonia till the excess of sulphydrate of ammonium is decomposed; acidify the liquid with hydrochloric acid, and then add a drop of solution of a ferric salt; a blood-red solution of sulphocyanide of iron will be formed.

This is a very delicate reaction. Some free sulphur in the yellow sulphydrate of ammonium unites wtth the alkaline cyanide and forms sulphocyanate ($2AmCy + S_2 = 2AmCyS$); the ammonia combines with excess of free sulphur and forms, among other salts, sulphydrate of ammonium, the whole of which is removed by the ebullition. If the liquid has not been evaporated far enough, sulphydrate of am-

*The Iodide of Hydrogen, or Hydriodic Acid*, is a heavy, colorless gas. Its solution in water may be made by passing sulphuretted hydrogen through water in which iodine is suspended.

$$2H_2S + 2I_2 = S_2 + 4HI.$$

Kolbe prepares hydriodic acid by adding to 10 parts of iodine contained in a flask, in an atmosphere of carbonic acid gas, 1 part of phosphorus, little by little and slowly. On the mixture of di- and tri-iodide of phosphorus are poured 4 parts of water. From this abundance of hydriodic acid is evolved on the application of gentle heat, and it is not contaminated by free iodine. Phosphoric acid remains.

*Iodide of potassium* (KI) is largely used in medicine, and hence is the most convenient iodide on which to experiment in studying the reactions of this acidulous radical. Solid iodine itself might be taken for the purpose; but its use and action in that state have already been alluded to in describing the iodides of potassium, cadmium, and mercury; its analytical reactions in the combined condition are those which may now occupy attention.

*Solution of Iodine.*—Iodine is slightly soluble in water (iodine-water), and readily soluble in an aqueous solution of iodide of potassium. Twenty grains of iodine and 30 of iodide of potassium dissolved in 1 ounce of distilled water, form *Liquor Iodi*, B. P.; *Liquor Iodinii Compositus*, U. S. P., is of the same character and about the same strength: 32 grains of iodine and 32 of iodide of potassium, rubbed with 1 fluidrachm of proof spirit, and 2 ounces of lard gradually mixed in, form *Unguentum Iodi*, B. P. *Unguentum Iodinii Compositum*, U. S. P., and *Unguentum Iodinii*, U. S. P., are similar preparations. It is more soluble in spirit (*Tinctura Iodinii*, U. S. P.), or in a spirituous solution of iodide of potassium (*Tinctura Iodi*, B. P., *Tinctura Iodinii Composita*, U. S. P.). It combines with sulphur, forming an unstable grayish-black solid iodide ($S_2I_2$), having a radiated crystalline structure (*Sulphuris Iodidum*, B. P. and U. S. P.; and *Unguentum Sulphuris Iodidi*). If 100 grains be thoroughly boiled with water, the iodine will pass off in vapor, and about 20 grains of sulphur remain.—*B. P.* and *U. S. P.*

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a few drops of an aqueous solution of an iodide (*e. g.* KI) add solution of nitrate of silver; a light yellow precipitate of iodide of silver (AgI) falls. Pour away the supernatant liquid and treat the precipitate with nitric acid, it is not dissolved; pour away the acid and then add dilute ammonia, it is only sparingly dissolved.

This reaction is useful in separating iodine from most other acidulous radicals, but does not distinguish iodine from bromine.

---

prismatic crystals. This, too, may have the formula $KI,I_2$. A mercuric hexiodide ($HgI_6$, perhaps $HgI_2,I_2,I_2$) is also known; and a periodide of ammonium, $NH_4I_2,I_2$.

Ammonia, it will be remembered, dissolves chloride of silver readily; hence the presence of chloride of potassium in bromide or iodide may be detected by dissolving in water, adding excess of nitrate of silver, collecting the precipitate, washing, digesting in ammonia, filtering, and adding excess of nitric acid to the filtrate; a white curdy precipitate indicates a chloride (of potassium). Bromide and iodide of silver are, however, slightly soluble in ammonia. A better process is given on page 276.

*Second Analytical Reaction.*—Liberate iodine from an iodide by the cautious addition of chlorine, then add cold decoction of starch; a deep-blue combination of iodine and starch, commonly termed "iodide of starch," is formed.

Starch is highly sensitive to the action of iodine; this reaction is consequently very delicate and characteristic. The reaction is not observed in hot liquids. Excess of chlorine must be avoided, or colorless chloride of iodine will be produced. Nitrous acid, or a nitrite acidulated with sulphuric acid, may be used instead of chlorine. Concentrated sulphuric acid also liberates iodine from iodides, the hydrogen of the hydriodic acid first produced uniting with the oxygen of the sulphuric acid—the latter ($H_2SO_4$) being reduced to sulphurous acid ($H_2SO_3$) or even to hydrosulphuric acid ($H_2S$).

In testing bromine for iodine the bromine must be nearly all removed by solution of sulphurous acid before the decoction of starch is added.

*Ozone* ($O_3$).—Papers soaked in mucilage of starch containing iodide of potassium form a test for free chlorine and nitrous acid, and are also employed by meteorologists to detect an allotropic or physically polymeric and energetic form of oxygen termed by Schönbein *ozone* (from ὀζω, *ozō*, I smell). This substance liberates iodine from iodide of potassium (with formation of iodide of starch), and is supposed to occur normally in the atmosphere, the salubrity or insalubrity of which is said to be dependent to some extent on the presence or absence of ozone. The possible occurrence of nitrous or chlorinoid gases in the air, however, renders the test untrustworthy. Houzeau proposes to test for ozone by exposing litmus paper of a neutral tint soaked in a dilute solution of iodide of potassium; the potash set free by action of the ozone turns the paper blue. The same paper without iodide would indicate the extent to which the effect might be due to ammonia vapor. Ozone, or rather ozonized air, is produced artificially in large quantities on passing air through a box (Beane's Ozone-generator) highly charged with electricity. In the latter operation condensation of the volume of air, or, rather, of the oxygen in the air, occurs. Small quantities are said to be obtained by exposing in a loosely closed bottle a stick of phosphorus partially covered by water, but the product is probably peroxide of hydrogen. It is a powerful bleaching, disinfecting, and general oxidizing agent; insoluble in water, soluble in oils of turpentine, cinnamon, and some other liquids. From experiments that have been made by Soret on the specific gravity of ozone, its molecular formula would seem to be

$O_3$, that of ordinary oxygen being $O_2$. Its smell is peculiar. (See p. 238, also "Blood.")

*Third Analytical Reaction.*—To a neutral aqueous solution of an iodide, add a solution containing one part of sulphate of copper to two and a half parts of green sulphate of iron, and well shake; a dirty-white precipitate of cuprous iodide ($Cu_2I_2$) falls.

$$2KI + 2CuSO_4 + 2FeSO_4 = Cu_2I_2 + K_2SO_4 + Fe_23SO_4.$$

Or to the liquid containing an iodide add the solution of copper sulphate and some solution of sulphurous acid, and warm the mixture, cuprous iodide falls.

$$2KI + 2CuSO_4 + H_2SO_3 + H_2O = Cu_2I_2 + 2KHSO_4 + H_2SO_4.$$

*Separation of Chlorides, Bromides, and Iodides.*—Chlorides and bromides are not affected in this way; the reaction is useful, therefore, in removing iodine from a solution in which chlorides and bromides have to be sought. The total removal of iodine by the former of the two modifications of the process is insured by supplementing the addition of the cupric and ferrous sulphates by a few drops of solution of potash or soda, any acid which might be keeping cuprous iodide in solution being thereby neutralized, ferric or ferrous hydrate, precipitated at the same time, not affecting the reaction. Occasionally, too, it may be necessary to repeat the process with the filtrate before the last traces of iodine are removed. The second modification of the process is, on the whole, to be preferred.

Chloride of the rare metal palladium performs a similar useful office in removing iodine, but not bromine or chlorine, from solutions.

Chlorides may be separated from bromides by taking advantage of the ready solubility of chloride of silver, and slow and slight solubility of bromide of silver in ammonia, especially in (a fair, not a great, excess of) ammonia containing chloride of silver.

Siebold's test for the detection of chlorides when occurring with bromides is based upon the fact that a strong solution of permanganate of potassium liberates the bromine from moderately strong solutions of bromides containing a large excess of sulphuric acid. A strong solution of permanganate is added to the aqueous solution of bromide or iodide (containing not more than 1 in 40), strongly acidified by dilute sulphuric acid, until the permanganate ceases to be decolorized, and a copious precipitate of oxide of manganese is formed. The mixture is boiled for about five minutes to expel the bromine, and then filtered. The colorless filtrate is now quite free from bromine, and may be tested for chlorine by nitrate of silver. If a chloride be present, a small quantity of its chlorine is lost by this process; but the main portion always remains, provided that no undue excess of the permanganate be used. It is essential, therefore, that the filtrate should be colorless; for if it be colored so as to indi-

cate the presence of undecomposed permanganate, the loss of the greater part if not of the whole of the chlorine must be expected. If, on the other hand, too little permanganate be employed, a trace of bromide may be left in the filtrate. If the solution under examination should be very much stronger than 1 in 40, water should be added before boiling (just after the addition of the permanganate), in order to avoid a loss of HCl.

Chlorides may also be detected in bromides and iodides by taking advantage of the formation of chlorochromic anhydride (page 239) and the non-occurrence of corresponding compounds of bromine or iodine.

To a solution of a chloride with a bromide and an iodide add a concentrated solution of sulphate of sodium, and then a reagent prepared by mixing equal volumes of sulphuric acid and saturated solution of sulphate of copper, until no further precipitation of cuprous iodide occurs. Next add solution of soda to remove excess of sulphate of copper; filter and evaporate to dryness. Place the dried residue, together with an equal bulk of red chromate of potassium, in a dry test-tube fitted with a delivery tube or into a small retort, and cover the mixture with sulphuric acid. Distil into water. Chromic anhydride and hydrochloric and hydrobromic acids are liberated by the sulphuric acid, and reacting upon one another form chlorochromic anhydride together with free bromine and chlorine.

$$CrO_3 + 2HCl = CrCl_2O_2 + H_2O.$$
$$2CrO_3 + 6HCl + 3H_2SO_4 = Cr_23SO_4 + 3Cl_2 + 6H_2O.$$
$$2CrO_3 + 6HBr + 3H_2SO_4 = Cr_23SO_4 + 3Br_2 + 6H_2O.$$

The chlorochromic anhydride is decomposed by the excess of water into which it falls, giving rise to chromic acid, which imparts its color to the liquid, and hydrochloric acid.

$$CrCl_2O_2 + 2H_2O = H_2CrO_4 + 2HCl.$$

The chlorine escapes and the bromine is dissolved by the water. The colored liquid is then shaken with chloroform, which removes the bromine—indicating bromine in the original substance. A yellow color remaining is due to chromic acid—indicating chlorides in the original substance. Or add ammonia to the distillate—the color due to bromine is thereby entirely removed while that of the chromic compound is only slightly modified.

*Fourth Analytical Reaction.*—Iodides have been shown to be useful in testing for mercuric salts (see the Mercury reactions, p. 196); a mercuric salt (corrosive sublimate, for example) may therefore be used in testing for iodides, a scarlet precipitate of mercuric iodide ($HgI_2$) being produced.

This reaction may be employed where large quantities of an iodide are present; but its usefulness in analysis is much impaired by the fact that the precipitate is soluble in excess of the dissolved iodide, or in excess of the mercuric reagent. Its color and insolubility in

with in commerce has a sp. gr. of 1.5, and contains 93 per cent. of real nitric acid ($HNO_3$); it fumes disagreeably, is unstable, and, except as an escharotic, is seldom used. The British and United States Pharmacopœias contain two acids: *Acidum Nitricum*, prepared as above, of sp. gr. 1.42 (also in U. S. P.), and containing 70 per cent. of real acid ($HNO_3$); and another, *Acidum Nitricum Dilutum*, sp. gr. 1.101 (U. S. P. 1.068), containing nearly 17¼ (17.44) per cent. Either of the stronger liquids, although containing water, is usually simply termed "nitric acid." The official nitric acid, of sp. gr. 1.42, is a definite hydrous acid ($2HNO_3,3H_2O$); it distils at 250° F. without change. If a weaker acid be heated it loses water, if a stronger acid be heated it loses nitric acid, until the density of 1.42 is reached. *Aqua fortis* is an old name for nitric acid (*Aqua fortis simplex*, sp. gr. 1.22 to 1.25; *Aqua fortis duplex*, 1.36). The strength of a specimen of nitric acid is determined by volumetric analysis. *Nitric anhydride* ($N_2O_5$), sometimes but erroneously called *anhydrous nitric acid*, is a solid crystalline substance formed on passing dry chlorine over dry nitrate of silver. Metals reduce nitric acid to nitrous acid and to the various oxides of nitrogen or even to nitrogen itself, according to the strength of acid, temperature, and amount of nitrate present. Not unfrequently nitrate of ammonium is simultaneously formed. Thus, with zinc:—

$$10HNO_3 + 2Zn_2 = 4(Zn2NO_3) + NH_4NO_3 + 3H_2O.$$

*Aqua Regia.*—Three fluidounces of nitric acid (B. P.), four of hydrochloric acid (B. P.), (3 to 5 by weight forms the *Acidum Nitromuriaticum*, U. S. P.), and twenty-five of water, give the *Acidum Nitrohydrochloricum Dilutum* of the British Pharmacopœia. The acids are ordered to be mixed twenty-four hours before dilution to insure mutual decomposition and full development of the chief active product, chlorine:—

$$\underset{\text{Nitric acid.}}{2HNO_3} + \underset{\text{Hydrochloric acid.}}{6HCl} = \underset{\text{Chloronitric gas.}}{N_2O_3Cl_4?} + \underset{\text{Water.}}{4H_2O} + \underset{\text{Chlorine.}}{Cl_2}$$

In the later stages of the reaction, the decomposition expressed in the following equation also probably occurs:—

$$\underset{\text{Nitric acid.}}{HNO_3} + \underset{\text{Hydrochloric acid.}}{3HCl} = \underset{\text{Chloronitrous gas.}}{NOCl} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Chlorine.}}{Cl_2}$$

The same reaction occurs if the acids are mixed after dilution, but is not complete for a week or a fortnight (Tilden). The undiluted mixture of acids is known as *aqua regia*, so called from its property of dissolving gold, "the king" of metals.

This "diluted nitrohydrochloric acid" is quite strong enough to attack organic matter with evolution of nitrous gases, hence should not be dispensed with tinctures, etc., in too concentrated a form.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a solution of any nitrate (*e. g.* $KNO_3$) add sulphuric acid, and then copper turnings, and

warm; colorless nitric oxide gas (NO) is evolved, which at once unites with the oxygen in the tube, giving *red fumes* of nitric peroxide or peroxide of nitrogen ($NO_2$).

$$2KNO_3 + 3H_2SO_4 + Cu_3 = 2NO + 3CuSO_4 + 4H_2O + 2KHSO_4;$$
$$2NO + O_2 = 2NO_2.$$

Performed on a larger scale, in a vessel to which a delivery-tube is attached, the reaction of nitric acid on copper becomes of synthetical interest, being the process for the preparation of nitric oxide gas for the purposes of chemical experiment.

Small amounts of a nitrate may be overlooked by this test, the color of the red fumes not being very intense.

Undiluted nitric acid poured on to copper turnings gives dense red vapors of *nitrous acid* ($HNO_2$), *nitrous anhydride* ($N_2O_3$), *nitric peroxide* ($NO_2$), *nitric oxide* (NO), and even *nitrogen* ($N_2$), the reaction varying somewhat according to the temperature of the mixture and (Ackworth) the amount of nitrate of copper in solution. Diluted nitric acid gives nitric oxide, $Cu_3 + 8HNO_3 = 3(Cu2NO_3) + 4H_2O + 2NO$.

*Second Analytical Reaction.*—To a cold solution of the nitrate, even if very dilute, add three or four crystals of sulphate of iron, shake gently for a minute in order that some of the sulphate may become dissolved, and then pour eight or ten drops of strong sulphuric acid down the side of the test-tube, so that it may form a layer at the bottom of the vessel; a reddish-purple or black coloration will appear between the acid and the supernatant liquid.

This is a very delicate test for the presence of nitrates. The black color is due to a solution or, perhaps, combination of nitric oxide with a portion of the ferrous salt. The nitric oxide is liberated from the nitrate by the reducing action of the hydrogen of the sulphuric acid, the sulphuric radical of which is absorbed by the ferrous sulphate, the latter salt becoming ferric sulphate.

$$2HNO_3 + 3H_2SO_4 + 6FeSO_4 = 4H_2O + 3(Fe_23SO_4) + 2NO.$$

*The process of oxidation* is one frequently employed in experimental chemistry; and nitrates, from their richness in oxygen, but more especially because always at hand, are the oxidizers usually selected for the purpose. In the operation they generally split up in one way, namely, into oxide of their basylous radical, nitric oxide gas, and available oxygen. Thus hydrogen nitrate (nitric acid) yields oxide of hydrogen (water) and the other bodies mentioned, as shown in the following equation:—

$$4NHO_3 = 2H_2O + 4NO + 3O_2.$$

When nitrates, other than nitric acid, are used for the purpose of oxidation, a stronger acid, generally sulphuric, is commonly added in

order that nitric acid may be formed; the hydrogen nitrate splitting up more readily than other nitrates.

*The five oxides of nitrogen* have now been mentioned, namely—

| | | | |
|---|---|---|---|
| Nitrous oxide . . . . | $N_2O$ | | $N_2O$ |
| Nitric oxide* . . . . | $NO$ | | $N_2O_2$ |
| Nitrous anhydride . . | $N_2O_3$ | or | $N_2O_3$ |
| Nitric peroxide* . . | $NO_2$ | | $N_2O_4$ |
| Nitric anhydride . . | $N_2O_5$ | | $N_2O_5$ |

Nitrous oxide is a colorless gas not altered by exposure to air; nitric oxide is also colorless, but gives red fumes in the air; nitrous anhydride is a red vapor condensible to a blue liquid; nitric peroxide is a red vapor condensible to an orange liquid; nitric anhydride is a colorless crystalline solid. The two anhydrides by absorbing water yield respectively nitrous acid ($HNO_2$) and nitric acid ($HNO_3$). Nitrous oxide is also probably an anhydride the acid of which would doubtless have the formula $HNO$, while the silver and sodium salts certainly have the formulæ $AgNO$ and $NaNO,3H_2O$ (Divers; Menke). The above series of compounds forms a good illustration of the doctrine of multiple proportions (p. 50).

*Third Analytical Reaction.*—Direct the blowpipe-flame on to charcoal until a spot is red-hot; now place on the spot a fragment of nitrate; deflagration ensues.

This reaction does not distinguish nitrates from chlorates. It is insufficient for the recognition of very small quantities of either class of salts, especially when they are mixed with other substances.

*Gunpowder* is an intimate mechanical mixture of 75 parts of nitre, 15 to 12½ parts of charcoal, and 10 to 12½ parts of sulphur. In burning it may be said to give sulphide of potassium (the white smoke), ($K_2S$), nitrogen ($N$), carbonic oxide ($CO$), and carbonic acid ($CO_2$) gases, though the decomposition is seldom complete. The sudden production of a large quantity of heated gas from a small quantity of a cold solid is sufficient to account for all the effects of gunpowder.

*Fourth Analytical Reaction.*—To nitric acid or other nitrate add solution of "sulphate of indigo;" the color is discharged.

"*Solution of Sulphate of Indigo*," B. P. (Sulphindylic or Sulphindigotic acid), is made by digesting 5 grains of dry finely powdered indigo in a small quantity of strong sulphuric acid in a test-tube for an hour, the mixture being kept hot by a water-bath; the blue liquid is then poured into 10 ounces of sulphuric acid, the whole well shaken, set aside, and the clear liquid decanted. Free chlorine also destroys the color of this reagent.

*Indigo*, B. P. ($C_8H_5NO$), is a blue coloring matter deposited when infusion of various species of *Indigofera* is exposed to air and slight warmth. Under these circumstances, *indican*, a yellow transparent

* The specific gravities of these gases indicate that $NO$ and $NO_2$ are the correct formulæ, and not $N_2O_2$ and $N_2O_4$.

amorphous substance, soluble in water, breaks up into indigo, which is insoluble and falls as a sediment, and a sort of sugar termed *indiglucin*. The indigo is collected, drained, pressed, and dried. By action of deoxidizing agents indigo is converted into soluble colorless *indigogen*, *reduced indigo* or *white indigo;* 1 part of powdered indigo, 2 of green sulphate of iron, 3 of slaked lime, and 200 of water, shaken together and set aside in a well-closed bottle, give this colorless indigo. A piece of yarn, calico, or similar fabric, dipped into such a solution, and exposed to air, becomes dyed blue, deposition of insoluble indigo-blue occurring within the cells and vessels of the fibre. This operation is readily performed on the small scale, and forms a good illustration of the characteristic feature of the art of dyeing, namely, the introduction of soluble coloring-matter into a fabric by permeation of the walls of its cellular and vascular tissue, and the imprisonment of that coloring-matter by conversion into a solid and insoluble form (*vide* also p. 139).

Pure indigo, or *indigotin*, may be obtained in beautiful needles by spreading a paste of indigo and plaster of Paris on a tin plate, and when dry placing a lamp underneath, moving the latter from place to place as the indigo sublimes and condenses on the surface of the plaster. It may also be obtained in crystals by gently boiling finely powdered indigo with aniline, filtering while hot, and setting aside; these crystals may be washed with alcohol. Hot paraffin may be employed instead of aniline. It is possible to obtain indigotin artificially.

*Distinction between nitric acid and other nitrates.*—Presence of the nitric radical in a solution having been proved by the above reactions, its occurrence as the nitrate of a metal is demonstrated by the neutral, or nearly neutral, deportment of the liquid with test-paper and the detection of the metal—its occurrence as nitric acid by the sourness of the liquor to the taste and the effervescence produced on the addition of a carbonate.

*Antidote.*—In cases of poisoning by strong nitric acid, solution of carbonate of sodium (common washing soda) or a mixture of magnesia and water may be administered as antidotes.

---

## QUESTIONS AND EXERCISES.

444. Trace the origin of nitrates.

445. In what does cubic nitre differ, chemically, from prismatic nitre?

446. Describe a process by which nitrate of potassium may be obtained artificially.

447. State the difference between nitrate of potassium, nitre, saltpetre, and sal prunella.

448. What group of elements is characteristic of all nitrates? and what claim has this group to the title of radical?

449. Mention the usual theory regarding the manner in which

atoms are arranged with reference to each other in such salts as nitrate of potassium.

450. How is Nitric Acid prepared?

451. Give the properties of nitric acid.

452. What reactions occur when strong nitric and hydrochloric acids are mixed?

453. How is nitric oxide prepared?

454. Enumerate and explain the tests for nitrates.

455. Into what substances does nitric acid usually split when employed as an oxidizing agent?

456. How is nitrous oxide prepared?

457. Enumerate the five oxides of nitrogen.

458. What is the nature of gunpowder?

459. Write a few sentences on the chemistry of indigo, one of the tests for nitric acid.

460. How is nitric acid distinguished from other nitrates?

461. What quantity of cubic nitre will be required to produce ten carboys of official nitric acid, each containing 114 pounds?—*Ans.* 1076⅔ pounds.

---

## CHLORIC ACID AND OTHER CHLORATES.

Formula of Chloric Acid $HClO_3$. Molecular weight 84.5.
Chlorates are made from hypochlorites.

### Hypochlorous Acid (HClO), and other Hypochlorites.

Place a few grains of red oxide of mercury in a test-tube, half-fill the tube with chlorine-water, and well shake the mixture; the resulting liquid is a solution of hydrochlorous acid, mercuric oxychloride remaining undissolved:—

$$2HgO + 2Cl_2 + H_2O = HClO + Hg_2OCl_2.$$

By the metathesis (double decomposition) of hydrochlorous acid and oxides or hydrates, other pure hypochlorites are formed:—

$$HClO + NaHO = NaClO + H_2O.$$

The direct action of chlorine on metallic hydrates and some carbonates is supposed to give a mixture of chloride and hypochlorite, as described in connection with the synthetical reactions of Sodium (p. 86, *Liquor Sodæ Chloratæ*, B. P.) and especially Calcium (p. 113, *Calx Chlorata*, B. P.).

$$Cl_2 + 2NaHO = NaCl,NaClO + H_2O;$$
$$2Cl_2 + 2CaH_2O_2 = CaCl_2,Ca2ClO + 2H_2O.$$

The condition of chlorides in these bodies is not satisfactorily made out; so that their constitution is not definitely deter-

mined. The action of acids on them results in the evolution of chlorine; hence the great value of the calcium compound (chlorinated lime, or chloride of lime) in bleaching operations:—

$$CaCl_2, Ca2ClO + 2H_2SO_4 = 2Cl_2 + 2CaSO_4 + 2H_2O.$$

The solubility of hypochlorites in water, their peculiar odor, greatly intensified on the addition of acid, and their bleaching-powers (see the above calcium reaction) are the characters on which to rely in searching for hypochlorites.

## Chlorates.

The group of elementary atoms represented by the formula $ClO_3$ is that characteristic of chloric acid and all other chlorates; hence it is expedient to regard it as being an acidulous radical, which may be termed the chloric radical. Like the nitric radical, it has not been isolated. Chloric anhydride ($Cl_2O_5$), unlike nitric anhydride, has not yet been obtained in the free condition.

Chlorates are artificial salts. They are formed by simply boiling aqueous solution of the common bleaching salts (chlorinated lime, chlorinated soda, chlorinated potash). Heat thus converts

| | | | | |
|---|---|---|---|---|
| 3(NaCl, NaClO) Chlorinated soda. | into | $NaClO_3$ Chlorate of sodium. | and | 5NaCl Chloride of sodium. |
| 3(KCl, KClO) Chlorinated potash. | into | $KClO_3$ Chlorate of potassium. | and | 5KCl Chloride of potassium. |
| 3($CaCl_2$, Ca2ClO) Chlorinated lime. | into | $Ca2ClO_3$ Chlorate of calcium. | and | 5$CaCl_2$ Chloride of calcium. |

One chlorate may also be made from another by double decomposition. In making chlorates economically the chlorinated salt is specially prepared for, and often in the same vessel as, the chlorate.

## Chlorate of Potassium.

Thus Chlorate of Potassium (*Potassii Chloras*, U. S. P.) is commercially made by saturating with chlorine gas a moistened mixture of three parts of chloride of potassium and ten of slaked lime, and well boiling the product. Chlorinated lime is first formed; this, on continued boiling with water, splits up into chloride of calcium and chlorate of calcium, and the latter reacting on the chloride of potassium yields chloride of calcium and chlorate of potassium.

$$6(Ca2HO) + 6Cl_2 = 3(CaCl_2, Ca2ClO) + 6H_2O;$$
$$3(CaCl_2, Ca2ClO) = Ca2ClO_3 + 5CaCl_2;$$
$$Ca2ClO_3 + 2KCl = CaCl_2 + 2KClO_3.$$

This operation may be conducted on a small scale by rubbing together in a mortar the above proportions of ingredients in ounces or half-ounces, adding enough water to make the whole assume the character of damp lumps, placing the porous mass in a funnel (loosely plugged with stones or pieces of glass), and passing chlorine gas (p. 27) through the mass by attaching the tube delivering the gas to the neck of the funnel. When the whole mass has become of a slight pink tint (due to a trace of permanganate), it should be turned into a dish, *well* boiled with water, filtered, the filtrate evaporated if necessary, and set aside; the chlorate of potassium crystallizes out in colorless rhomboidal plates, chloride of calcium remaining in the mother-liquor.

In the official process, carbonate of potassium instead of chloride is used; but otherwise it is similar to the method just described. Chlorinated potash and chlorinated lime are first formed:—

$$K_2CO_3 + Ca2HO + Cl_2 = KCl,KClO + CaCO_3 + H_2O,$$
$$6(Ca2HO) + 6Cl_2 = 3(CaCl_2,Ca2ClO) + 6H_2O;$$

these, on boiling with water, split up into chlorates and chlorides—

$$3(KCl,KClO) = KClO_3 + 5KCl,$$
$$3(CaCl_2,Ca2ClO) = Ca2ClO_3 + 5CaCl_2;$$

the whole of the chloride of potassium and chlorate of calcium finally yielding chlorate of potassium and chloride of calcium—

$$2KCl + Ca2ClO_3 = CaCl_2 + 2KClO_3.$$

Neglecting intermediate decomposition, the reactions may be represented by the following equation:—

$$\underset{\text{Chlorine.}}{6Cl_2} + \underset{\text{Carbonate of potassium.}}{K_2CO_3} + \underset{\text{Hydrate of calcium.}}{6CaH_2O_2} = \underset{\text{Chlorate of potassium.}}{2KClO_3} + \underset{\text{Carbonate of calcium.}}{CaCO_3} + \underset{\text{Chloride of calcium.}}{5CaCl_2} + \underset{\text{Water.}}{6H_2O.}$$

Chlorate of potassium is soluble in water to the extent of 6 or 7 parts in 100 at common temperatures. It is usually administered medicinally in aqueous solution, sometimes also in lozenges (*Trochischi Potassii Chloratis*, U. S. P.). Chlorate of potassium must on no account be rubbed with sulphur in a mortar or otherwise, friction of such a mixture resulting in violent explosion.

Chlorate of potassium, when heated, yields chloride of potassium and oxygen, and is the salt commonly employed in the preparation of the gas for experimental purposes. But if the action be arrested when one-third of the oxygen has escaped, the residual salt is found to contain perchlorate of potassium ($KClO_4$):—

$$2KClO_3 = KClO_4 + KCl + O_2.$$

*Perchloric Acid* ($HClO_4$).—Crude perchlorate of potassium, obtained as just indicated, is boiled (in a fume-cupboard) with hydro-

chloric acid to decompose any chlorate that may be remaining, and then separated from chloride by washing and crystallization, chloride being far more soluble in water than the perchlorate. Perchloric acid is then obtained by distilling the perchlorate of potassium with sulphuric acid; it is quite stable, and is occasionally administered in medicine.

*Chloric Acid* ($HClO_3$) may be isolated, but is unstable, quickly decomposing into chlorine, oxygen, and perchloric acid; some other chlorate (*e. g.*, $KClO_3$) must therefore be used in studying the reactions of the chloric radical.

*Table of the Chlorine Acids.*

| | |
|---|---|
| Hydrochloric acid . . . . | HCl. |
| Hypochlorous acid . . . . | HClO. |
| Chlorous acid . . . . . | $HClO_2$. |
| Chloric acid . . . . . | $HClO_3$. |
| Perchloric acid . . . . . | $HClO_4$. |

*The chloric radical is univalent* ($ClO_3$). The acidulous radicals of the other chlorine acids are also univalent, as indicated in the foregoing formulæ.

## *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To solution of a chlorate (*e. g.*, chlorate of potassium) add solution of nitrate of silver; no precipitate falls, showing that the chlorine must be performing different functions from those it possesses in chlorides. Evaporate the solution to dryness, and place the residue in a small dry test-tube, or at once drop a fragment of a chlorate into a test-tube, and heat strongly; oxygen is evolved, and may be recognized by its power of reinflaming an incandescent match inserted in the tube. Boil the residue with water, and again add solution of nitrate of silver; a white precipitate falls, having all the characters of chloride of silver, as described under hydrochloric acid.

This is a trustworthy test, and, even omitting the recognition of the oxygen, may be applied in the detection of small quantities of chlorates.

*Second Analytical Reaction.*—To a fragment of a chlorate add two or three drops of strong sulphuric acid; an explosive gas ($Cl_2O_4$) is evolved somewhat resembling chlorine in odor but possessing a deeper color than that element.

$$3KClO_3 + H_2SO_4 = Cl_2O_4 + KClO_4 + K_2SO_4 + H_2O.$$

Warm the upper part of the test-tube to 150° or 200° F., or introduce a hot wire; a sharp explosion ensues, due to decomposition of the gas, peroxide of chlorine, into its elements.

*Third Analytical Reaction.*—Heat a small fragment of a chlorate with hydrochloric acid: a yellowish-green explosive gas termed *euchlorine* is evolved. Its color is deeper than that of chlorine, hence the name (from εὖ, *eu*, well, and χλωρός, *chlōros*, green). In odor it resembles chlorine, and is probably a mixture of that element with one of the oxides of chlorine.

*Fourth Analytical Reaction.*—Direct the blowpipe-flame on to charcoal until a spot is red-hot, and then place on the spot a fragment of a chlorate; deflagration ensues as with nitrates.

## Bromates.

Bromates are salts closely resembling chlorates and iodates. The formula of Bromic Acid is $HBrO_3$.

## Iodates.

*Iodic Acid* ($HIO_3$).—Iodine is boiled in a flask with five times its weight of the strongest nitric acid (sp. gr. 1.5), in a fume-cupboard, until all action ceases. On cooling iodic acid separates in small pyramidal crystals. These are removed, the residual liquid evaporated to dryness to remove excess of nitric acid, the residue and the first crop dissolved in a small quantity of boiling water, and the solution set aside to crystallize.

*Iodide of Potassium* ($KIO_3$).—Powder together equal weights of iodine and chlorate of potassium; to the mixture add twice its weight of water and about one-eighth of its weight of nitric acid; warm the whole until iodine disappears, and evaporate quite to dryness over a water-bath. The residue dissolved in water forms "Solution of Iodate of Potash," B. P. It contains a little nitrate of potassium.

In this reaction the small quantity of nitric acid furnishes corresponding amounts of nitrate of potassium and chloric acid. The chloric acid with iodine gives iodic acid and chlorine, thus:—

$$2HClO_3 + I_2 = 2HIO_3 + Cl_2.$$

The iodic acid and some chlorate of potassium then yield chloric acid and iodate of potassium—

$$HIO_3 + KClO_3 = HClO_3 + KIO_3;$$

and the two reactions alternate until the whole of the iodine has displaced the whole of the chlorine.

Iodate of potassium and sulphurous acid decompose each other with elimination of iodine (or with formation of a blue color, if starch be present). Sulphurous acid occurring as an impurity in acetic and other acids may thus be detected.

$$2KIO_3 + 5H_2SO_3 = I_2 + 3H_2SO_4 + 2KHSO_4 + H_2O.$$

*Ferric iodate*, or rather *Oxyiodate* ($Fe_2O4IO_3,8H_2O$), is precipitated on adding solution of ferric chloride to solution of iodate of potassium.

## QUESTIONS AND EXERCISES.

462. How may hypochlorous acids be formed?
463. What are the relations of hypochlorous acid to common bleaching powder?
464. By what reaction is chlorine eliminated from hypochlorites?
465. State the general reaction by which chlorates are formed.
466. Give details of the preparation of chlorate of potassium.
467. Mention the properties of chlorate of potassium.
468. What decompositions occur when chlorate of potassium is heated?
469. Find the molecular weight of chlorate of potassium.
470. What weight of oxygen is yielded when 1 oz. of chlorate of potassium is completely decomposed, and how much chloride of potassium remains?
471. One hundred cubic inches of oxygen, at 60° F. and barometer at 30 inches, weighing 34.203 grains, and 1 gallon containing 277¼ cubic inches, what weight of chlorate of potassium will be required to yield 10 gallons of the gas? Ans. 5½ ounces.
472. How many cubic inches of oxygen are producible from 1 oz. of chlorate of potassium?
473. Calculate the weight of chlorate of potassium theoretically obtainable from 100 parts of chloride.
474. How is perchloric acid prepared?
475. Enumerate the chlorine acids.
476. How may the presence of chlorides in chlorates be demonstrated?
477. Mention the tests for chlorates.
478. Give the formula of peroxide of chlorine.
479. What is euchlorine?
480. How may iodic acid be made?
481. Describe the preparation of iodate of potassium.

---

## ACETIC ACID AND OTHER ACETATES.

Formula of Acetic Acid $HC_2H_3O_2$, or $H\overline{A}$. Molecular weight 60.

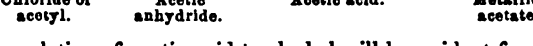
*Source.*—Acetic acid is said to occur naturally in certain plant juices and animal fluids in minute proportions, but otherwise is an artificial product. Much is furnished by the destructive distillation of wood, hence the term *pyroligneous acid* for the crude product, a hybrid word from πῦρ, *pūr*, fire, and *lignum*, wood. This impure product, neutralized by carbonate of sodium, the whole evaporated and the residue gently heated to drive off the volatile tarry matters, gives acetate of sodium, which after recrystallization furnishes by distillation with oil of vitriol and water acetic acid in a fair state of purity. In Germany and France large quantities of acetic acid are made by the spontaneous oxidation of the alcohol in inferior wines, hence the *white-* and *red-wine vinegar* (*vinegar*, from the French

*vin*, wine, and *aigre*, sour). In England also the domestic form of acetic acid (brown vinegar) has a similar origin: infusion of malt and unmalted grain is fermented; and the resulting oxidation of its sugar, instead of being arrested when the product is an alcoholic liquid, a sort of beer, is allowed to go on to the next stage, acetic acid; it usually contains from 3 to 6 per cent. of real acetic acid ($HC_2H_3O_2$).

*Vinegars.*—The official vinegar (*Acetum*, U. S. P.) contains 5½ (5.4) per cent. of acid. The so-called Vinegar of Cantharides (*Acetum Cantharidis*, B. P.) is a solution of the active principle of cantharides in very strong acetic acid, not in vinegar. The Vinegar of Squill (*Acetum Scillæ*, B. P. and U. S. P.) is also a solution of the active principle of squill in dilute acetic acid, not in true vinegar. The same may be said of *Acetum Lobeliæ*, U. S. P., *Acetum Opii*, U. S. P., and *Acetum Sanguinariæ*, U. S. P. (Vinegar of Bloodroot). Distilled vinegar (*Acetum Destillatum*, U. S. P.) is a form of Dilute Acetic Acid not now official in Great Britain. The *Acetum Opii* or *Black Drop* of America is made from nutmegs, saffron, and sugar, as well as Opium and Diluted Acetic Acid. In the British Pharmacopœia, vinegar, except its use for its own sake, is only employed in the preparation of *Emplastrum Cerati Saponis*.

*The Acetic Radical.*—The group of elements represented by the formula $C_2H_3O_2$ is that characteristic of acetic acid and other acetates, and may, for convenience of study, be assumed to be an acidulous univalent radical. It has not been isolated, unless indeed a compound of similar composition, resulting from the action of peroxide of barium on acetic anhydride, is the radical in question.

*Acetyl.*—The characteristic grouping in acetates, $C_2H_3O_2$, is frequently considered to contain, rather than to be, a radical—$C_2H_3O$, termed *acetyl*. Acetates may be made to yield a body having the composition $C_2H_3OCl$, which is regarded as chloride of acetyl; from this may be obtained acetic anhydride ($C_4H_6O_3$), which by absorbing water becomes acetic acid.




The relation of acetic acid to alcohol will be evident from the following equation representing empirically the formation of the acid:—

$$\underset{\text{Alcohol.}}{C_2H_6O} + O_2 = \underset{\text{Acetic acid.}}{C_2H_4O_2} + H_2O.$$

*Acetates* in aqueous solution are liable to decomposition. In solution of acetate of morphia a myceloid growth occasionally forms, acetic acid disappears, and morphia is deposited. Solution of acetate of ammonium is liable to a similar change, gradually becoming alkaline.

*Synthetical Reaction.*

## Acetic Acid.

*Synthetical Reaction.*—To a few grains of acetate of sodium in a test-tube add a little water and some sulphuric acid, and heat the mixture; acetic acid is evolved, and may be condensed by a bent tube adapted to the test-tube by a cork in the usual way.

*Acetic Acid.*—This is the process by which acetate of sodium or calcium (the neutralized products of the distillation of wood) is made to yield acetic acid on the large scale. As with nitric and hydrochloric acids, the loose term "acetic acid" is that usually applied to aqueous solutions of acetic acid. The *Acidum Aceticum*, B. P., contains nearly 33 per cent. of real acid, that is, of $HC_2H_3O_2$; for it contains only 28 per cent. of *acetic anhydride* ($C_4H_6O_3$)—still occasionally though somewhat obscurely termed *anhydrous acetic acid.* Its specific gravity is 1.044, the specific gravity of *Acidum Aceticum*, U. S. P., being 1.047. *Acidum Aceticum Dilutum*, B. P. and U. S. P., contains 4½ per cent. of $HC_2H_3O_2$. Glacial acetic acid ($HC_2H_3O_2$) contains no water. It solidifies to a crystalline mass at temperatures below 63° F., hence the appellation *glacial* (from *glacies*, ice.) Good commercial glacial acetic acid (*Acidum Aceticum Glaciale*, B. P.) does not contain more than 1 per cent. of water, corresponding to 84.15 per cent. of acetic anhydride; it solidifies at 34° F. and again liquefies at 48°: its specific gravity is 1.065. Although water is lighter than this acetic acid, yet the addition of water at first renders the acid heavier; evidently therefore condensation, or contraction in bulk, occurs on mixing the liquids: after 10 per cent. has been added, the addition of more water produces the usual effect of dilution of a heavy liquid by a lighter, namely, reduction of relative weight. This matter will be better understood after the subject of specific gravity has been studied.

The following equation is expressive of the above reaction:—

$$\underset{\text{Acetate of sodium.}}{NaC_2H_3O_2} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Acetic acid.}}{HC_2H_3O_2} + \underset{\text{Acid sulphate of sodium.}}{NaHSO_4}$$

or, assuming the existence of acetyl ($C_2H_3O$) in acetic acid, and a corresponding radical sulphuryl ($SO_2$) in sulphuric acid—

$$\left.\begin{matrix}C_2H_3O\\Na\end{matrix}\right\}O + \left.\begin{matrix}SO_2\\H_2\end{matrix}\right\}O_2 = \left.\begin{matrix}C_2H_3O\\H\end{matrix}\right\}O + \left.\begin{matrix}SO_2\\NaH\end{matrix}\right\}O_2;$$

or, thirdly, on the assumption that salts contain the oxide of a basylous radical united with the anhydride of an acid (the old view under which such names as acetate of soda were formed)—

$$Na_2O,C_4H_6O_3 + 2H_2O,SO_3 = Na_2O,H_2O,2SO_3 + H_2O,C_4H_6O_3.$$

*Note on the Constitution of Salts.*

Which of these three equations, or, more broadly, which of the three views of the constitution of salts illustrated by the equations, is correct, it is impossible to say. Whether it is $C_2H_3O_2$, $C_2H_3O$, or $C_4H_6O_3$, which migrates from one acetic compound to another, whether it is $SO_4$, $SO_3$, or $SO_2$, which migrates from one sulphuric compound to another, and so on with other acidulous groupings, cannot at present be determined. There are strong objections to each view; and possibly neither is right. Either the given radicals cannot be isolated, or application of the forces of heat, light, and electricity do not confirm views arrived at by the results of operations with the chemical force; or a salt comes to be regarded as having so large a number of constituent parts that the view, however true, breaks down in practice from the sheer inability of the mind to grasp the complicated analogies involved. Yet for the purposes of description, study, and conversation some system must be adopted. Let the first, then, be generally taken, over-reliance on it being checked by the use of general instead of special names for the hypothetical radicals, and other systems be employed in certain cases. (See also p. 286.)

*Impurity.* — Acetic acid often contains sulphurous acid. The methods by which this impurity is detected are described on page 307.

*Analytical Reactions (Tests).*

*First Analytical Reaction.*—To an acetate add sulphuric acid and heat the mixture; acetic acid, recognized by its odor, is evolved.

*Note* 1.—Iodine, sulphurous acid, and other substances of powerful odor, mask that of acetic acid; they must be removed, therefore, usually by precipitation or oxidation, before applying this test.

*Note* 2. It will be noticed that this reaction is identical with the previous one; it has synthetical or analytical interest, according to the object and method of its performance.

*Second Analytical Reaction.*—Repeat the above action, a few drops of spirit of wine being first added to the acetate; acetic ether (acetate of ethyl, $C_2H_5C_2H_3O_2$), also of characteristic odor, is evolved.

The basylous radical ethyl ($C_2H_5$) will be referred to subsequently.

*Third Analytical Reaction.*—Heat a fragment of a dry acetate in a test-tube, and again notice the odor of the gaseous products of the decomposition; among them is *acetone* ($C_3H_6O$), the smell of which is characteristic. Carbonate of the metal remains in the test-tube.

*Fourth Analytical Reaction.*—To a solution of an acetate made neutral by the addition of acid or alkali, as the case may

be, add a few drops of neutral solution of perchloride of iron; a deep-red liquid results, owing to the formation of ferric acetate ($Fe_2 6C_2H_3O_2$).

*Analytical Note.*—It will be noticed that the formation of characteristic precipitates, the usual method of removing radicals from solution for recognition, is not carried out in the qualitative analysis of acetates. This is because all acetates are soluble. Acetate of silver ($AgC_2H_3O_2$) and mercurous acetate ($HgC_2H_3O_2$) are only sparingly soluble in cold water, but the fact can seldom be utilized in analysis. Hence peculiarities of color and odor, the next best characters on which to rely, are adopted as means by which acetates may be detected. Acetates, like other organic compounds, char when heated to a high temperature.

*Note on Anhydrides.*—Up to this point the student has regarded an anhydride as a body derived from an acid by removal of the *whole* of the hydrogen of the acid, together with as much of its oxygen as with the hydrogen forms water. The definition will scarcely apply to acetic anhydride, and must therefore be somewhat qualified. An *anhydride* is derived from an acid, the acid having lost the whole of its *basylous* hydrogen and so much oxygen as is necessary to form water with that hydrogen. Anhydrides are obtained by heating acids, and by other methods.

## QUESTIONS AND EXERCISES.

482. What is the formula of acetic acid?
483. State the relation of acetic acid to other acetates.
484. What is the molecular weight of acetic acid?
485. Name the sources of acetic acid.
486. What is pyroligneous acid?
487. From what compound is the acetic acid of foreign and English vinegar immediately derived?
488. How much real acid is contained in official vinegar?
489. What is the nature of the "Vinegars" of Pharmacy?
490. How may acetic acid be obtained from acetate of sodium?
491. How much real acid is contained in the official acetic acid?
492. Mention the strength of commercial glacial acetic acid?
493. Give three or more views of the constitution of acetates, illustrating each by formulæ.
494. Enumerate the tests for acetates.

## HYDROSULPHURIC ACID AND OTHER SULPHIDES.

Formula of Hydrosulphuric Acid $H_2S$. Molecular weight 34.

*Source and Varieties of Sulphur.*—The acidulous radical of *hydrosulphuric acid*, *sulphydric acid*, or *sulphuretted hydrogen*, and other sulphides, is the element sulphur (S). It occurs in nature in

combination with metals, as already stated in describing the ores of some of the metals, and also in the free state. Most of the sulphur used in medicine is imported from Sicily, where it occurs chiefly associated with blue clay. It is purified by fusion, sublimation, or distillation. Melted and poured into moulds, it constitutes a crystalline mass termed *roll sulphur*. If distilled and the vapor carried into large chambers, so that it may be rapidly condensed, the crystals are so minute as to give the sulphur a pulverulent character; this is *sublimed sulphur* (*Sulphur Sublimatum*, B. P. and U. S. P.) or *flowers of sulphur:* the same washed constitutes *Sulphur Lotum*, U. S. P.

The third common form, *milk of sulphur*, will be noticed subsequently. Sulphur also occurs in nature in combination as a constituent of animal and vegetable tissues, as sulphurous acid gas ($SO_2$) in volcanic vapors, and as sulphuretted hydrogen in some waters, as those of Harrogate.

*Plastic sulphur* is one of the allotropic varieties of the element, obtained on heating sulphur considerably beyond its melting-point and pouring into cold water.

*Quantivalence.*—Sulphur is sexivalent, as seen in sulphuric anhydride ($SO_3$), a substance which will be noticed under sulphuric acid. It also occasionally exhibits quadrivalent ($SO_2$) and still oftener bivalent affinities ($H_2S$).

*Molecular weight.*—At very high temperatures sulphur follows the rule that, under similar conditions of heat and pressure, atomic weights (in grammes, grains, etc.) of volatile elements occupy equal volumes of vapor; its formula therefore is $S_2$, and molecular weight 64. At lower temperatures the volume weighs three times as much as it should do if following usual laws, and then the molecule would appear to contain six atoms ($S_6$).

*Acid Salts.*—Sulphur ($S''$) being the first acidulous radical of bivalent activity met with in these sections on acids, it is desirable here to draw attention to a new class of salts to which such a radical will generally give rise. These are acid salts, which are intermediate between normal salts and acids. Univalent radicals with an atom of hydrogen give an acid, and with an atom of other basylous radicals an ordinary or normal salt. But bivalent radicals, from the fact that they give with two atoms of hydrogen an acid, and with two atoms of univalent metals a normal salt, may obviously give intermediate bodies containing one atom of hydrogen and one atom of metal: these are appropriately termed acid salts: they are neither normal acids nor normal salts, but *acid salts*. (Examples:—$KHCO_3$, $NaHSO_4$, $KHC_4H_4O_6$, $Na_2HPO_4$, $CuHAsO_3$, $CaH_42PO_4$.) Whether or not these salts give an acid reaction with blue litmus paper depends on the strength of the respective radicals. Usually they do redden the test paper, but sometimes not; thus the acid sulphide or sulphydrate of potassium (KHS), of sodium (NaHS), or ammonium (AmHS) has alkaline properties.*

---

* Some chemists regard these sulphydrates as compounds of basylous radicals with HS, a univalent grouping termed hydrosulphyl (persulphide of hydrogen, $H_2S_2$), just as hydrates are similarly viewed

*Synthetical Reactions.*

## Sulphuretted Hydrogen.

*First Synthetical Reaction.*—The preparation of sulphuretted hydrogen.—This operation was described on page 96, and probably has already been studied by the reader.

## Precipitated Sulphur.

*Second Synthetical Reaction.*—Prepare the variety of the radical of sulphides known as *Precipitated Sulphur* (*Sulphur Præcipitatum*, B. P. and U. S. P.), or *milk* of sulphur, by boiling a few grains of flowers of sulphur (5 parts) with slaked lime (3 parts) (10 parts U. S. P.) and some water (20 parts) in a test-tube (larger quantities in an evaporating-basin), filtering, and (reserving a small portion of the filtrate) adding dilute hydrochloric acid until the well-stirred milk-like liquid has a faint acid reaction on test-paper; sulphur is precipitated, and may be collected on a filter, washed, and dried (at about 120°). Excess of acid must be avoided, or some hydrosulphyl, the liquid persulphide of hydrogen ($H_2S_2$) will be formed, probably causing the particles of sulphur to aggregate to a gummy mass.

This is the process of the Pharmacopœias. Polysulphide of calcium and hyposulphite of calcium are formed:—

$$\underset{\text{Hydrate of calcium.}}{3CaH_2O_2} + \underset{\text{Sulphur.}}{6S_2} = \underset{\text{Polysulphide of calcium.}}{2CaS_5} + \underset{\text{Hyposulphite of calcium.}}{CaS_2O_3} + \underset{\text{Water.}}{3H_2O}$$

On adding the acid, both salts are decomposed and sulphur separates:—

$$\underset{\text{Polysulphide of calcium.}}{2CaS_5} + \underset{\text{Hyposulphite of calcium.}}{CaS_2O_3} + \underset{\text{Hydrochloric acid}}{6HCl} = \underset{\text{Chloride of calcium.}}{3CaCl_2} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Sulphur.}}{6S}$$

The polysulphide of calcium yields sulphuretted hydrogen as well as milky-white sulphur on the addition of acid. The hyposulphite of calcium then yields sulphurous acid gas as well as a yellowish sulphur.

---

as compounds of the univalent radical hydroxyl (HO) (peroxide of hydrogen, $H_2O_2$)—$H_2S$ becoming HHS or HIIs (hydrosulphylide of hydrogen), and $H_2O$ becoming HHO or HHo (hydroxylide of hydrogen). The chemical analogy between sulphur and oxygen, already once alluded to (p. 176), is further illustrated by the compound just mentioned. Sulphur is also closely related to the rarer element Selenium. Thus we have $SeO_2$ as well as $SO_2$, $H_2SeO_3$ (selenious acid) as well as $H_2SO_3$ (sulphurous acid), $H_2SeO_4$ (selenic acid) as well as $H_2SO_4$ (sulphuric acid). The rare element Tellurium also seems to have similar analogies. The four hydrogen compounds of the group have the formulæ $H_2O$, $H_2S$, $H_2Se$, $H_2Te$.

These gases react and give sulphur and water, very little sulphuretted hydrogen escaping. A little pentathionic acid (see Index) is also said to be formed.

$$4H_2S + 2SO_2 = 3S_2 + 4H_2O.$$

*Impure Precipitated Sulphur.* The *old* "Milk of Sulphur."—To a sulphur solution prepared as before (or to the reserved portion) add a little dilute sulphuric acid; the precipitate is in this case largely mixed with sulphate of calcium:—

$$\underset{\text{Poly-sulphide of calcium.}}{2CaS_5} + \underset{\text{Hyposulphite of calcium.}}{CaS_2O_3} + \underset{\text{Sulphuric acid.}}{3H_2SO_4} = \underset{\text{Sulphate of calcium.}}{3CaSO_4} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Sulphur.}}{6S_2}$$

Place a little of each of these specimens of precipitated sulphur with a drop of the supernatant liquid on a strip of glass, cover each spot with a piece of thin glass, and examine the precipitates under a microscope; the pure sulphur will be found to consist of minute grains or globules, the impure to contain comparatively large crystals (sulphate of calcium).

*Note.*—A large proportion of the precipitated sulphur met with in trade in England still is thus mixed with sulphate of calcium, most specimens containing two-thirds of their weight of that substance. Many purchasers, indeed, are so accustomed to the satiny appearance of the mixed article as to regard the real sulphur with suspicion, sometimes refusing to purchase it. The mixed article is, certainly, more easily mixable with aqueous liquids; but it has long been discarded from the Pharmacopœias of Great Britain, and, on the whole, is scarcely worthy of a prominent place in the pharmacy of 1879.

Many pharmacists now refuse all parcels of sulphur which yield a white ash (the anhydrous sulphate) when a little is burnt off on the end of a table-knife or spatula. (No more damage is done to the steel than a rub on a knife-board will remove.)

To ascertain exactly the amount of sulphate of calcium in a specimen of impure precipitated sulphur, place a weighed quantity in a tared crucible and heat till no more vapors are evolved. The weight of the residual anhydrous sulphate of calcium ($CaSO_4 = 136$), with one-fourth thereof added, is the amount of crystalline sulphate of calcium ($CaSO_4,2H_2O = 172$) present in the original quantity of impure sulphur.

### *Analytical Reactions (Tests).*

To a sulphide add a few drops of hydrochloric acid; sulphuretted hydrogen will probably be evolved, well known by its smell. If the sulphide is not acted upon by the acid, or if free sulphur be under examination, mix a minute portion with a fragment of solid caustic potash or soda, and fuse on a silver coin or spoon. When cold, place a drop of dilute hydrochloric

acid on the spot, sulphuretted hydrogen is evolved, and a black stain, due to sulphide of silver ($Ag_2S$), left on the coin.

Other sulphur reactions may be adopted as tests, but the above are sufficient for all ordinary purposes. The most convenient reagent for detecting a sulphide in solution of ammonia is ammonio-sulphate of copper, which gives a black precipitate of sulphide of copper if a sulphide be present.

The *Iodide of Sulphur* ($S_2I_2$) has been mentioned under "Iodine." A chloride ($S_2Cl_2$) and bromide ($S_2Br_2$) may also be formed from their elements. A mixture of sulphur and chloride of sulphur is sometimes met with under the name of *hypochloride of sulphur*.

## QUESTIONS AND EXERCISES.

495. In what form does sulphur occur in nature?

496. State the modes of preparation of the three chief commercial varieties of sulphur.

497. To what extent does the atom of sulphur vary in quantivalence?

498. State the relations of acid salts to acids and to normal salts.

499. Define sulphides and sulphydrates.

500. Describe the preparation of sulphuretted hydrogen.

501. What are the characters of pure precipitated sulphur?

502. Give equations explanatory of the reactions which occur in precipitating sulphur according to the official process.

503. Describe the microscopic test for impure precipitated sulphur.

504. Mention a ready physical method of detecting sulphate of calcium in precipitated sulphur.

505. Mention the tests for sulphides, and the character by which sulphuretted hydrogen is distinguished from other sulphides.

506. How are sulphides insoluble in acids tested for sulphur?

507. Give a method for the detection of a trace of sulphur in solution of ammonia.

## SULPHUROUS ACID AND OTHER SULPHITES.

Formula of sulphurous acid $H_2SO_3$. Formula of sulphurous acid gas or sulphurous anhydride, commonly termed sulphurous acid, $SO_2$. Molecular weight of sulphurous acid 82; of the anhydride 64.

When sulphur is burned in the air it combines with oxygen and forms sulphurous acid gas ($SO_2$), more correctly termed sulphurous anhydride, or commonly, but erroneously, sulphurous acid. It is a pungent, colorless gas, readily liquefied on being passed through a tube externally cooled by a *freezing-mixture* composed of two parts of well-powdered ice (or, better, snow) with one part of common salt. If sulphurous acid gas becomes moist or is passed into water, heat is

evolved and true sulphurous acid ($H_2SO_3$) formed. The latter body may be obtained in crystals; but it is very unstable, and hence the properties of the sulphurous radical must be studied under the form of some other sulphite; sulphite of calcium ($CaSO_3$) or sulphite of sodium ($Na_2SO_3$) may be used for the purpose.

*Quantivalence.*—The radical of the sulphite is bivalent ($SO_3''$), and hence forms acid sulphites, such as acid sulphite of potassium ($KHSO_3$), and normal sulphites, such as sulphite of sodium ($Na_2SO_3$).

*Note on Nomenclature.*—The sulphites are so named from the usual rule, that salts corresponding with acids whose names end in *ous* have a name ending in *ite*. They are generally made by passing sulphurous acid gas over moist oxides or carbonates—in the latter case carbonic acid gas escapes.

*Synthetical Reaction.*—To a few drops of sulphuric acid in a test-tube add a piece of charcoal and apply heat; sulphurous acid gas is evolved, and may be conveyed by a bent tube into a small quantity of cold water in another test-tube. Larger quantities may be made in a Florence flask. The product is the *Acidum Sulphurosum*, B. P. and U. S. P. It is said to contain, if saturated, nearly 12 (11.79) per cent. of sulphurous acid ($H_2SO_3$) or about 9 (9.2) per cent. of the gas ($SO_2$). The process is also that described in the Pharmacopœia, except that the gas is purified by passing through a small wash-bottle before final collection. Specific gravity 1.04 (1.035, U. S. P.).

$$\underset{\text{Sulphuric acid.}}{4H_2SO_4} + \underset{\substack{\text{Carbon}\\\text{(charcoal).}}}{C_2} = \underset{\text{Carbonic acid gas.}}{2CO_2} + \underset{\text{Water.}}{4H_2O} + \underset{\text{Sulphurous acid gas.}}{4SO_2}$$

$$\underset{\text{Sulphurous acid gas.}}{SO_2} + \underset{\text{Water.}}{H_2O} = \underset{\text{Sulphurous acid.}}{H_2SO_3}$$

Sulphurous acid gas may also be made by boiling copper, mercury, or iron with sulphuric acid, sulphates of the metals being formed. Also by boiling sulphur with sulphuric acid.

If in this process the water were replaced by solutions of or solid metallic oxides or carbonates, sulphites of the various metals would be formed. The formula of sulphite of potassium (*Potassii Sulphis*, U. S. P.) is $K_2SO_3,2H_2O$; of sulphite of sodium (*Sodii Sulphis*, U. S. P.) $Na_2SO_3,7H_2O$; of the bisulphite or acid sulphite $NaHSO_3$. Under the name of *antichlor* the former is used for removing traces of chlorine from paper pulp. The so-called *Bisulphite of Lime*, used by brewers for retarding or arresting fermentation and oxidation, and for various antiseptic purposes, is a solution of sulphite of calcium ($CaSO_3$) in free sulphurous acid ($H_2SO_3$), and is made by passing sulphurous acid gas ($SO_2$) into thin milk of lime. Its specific gravity varies from 1050 to 1070, and its potential strength of anhydride ($SO_2$) from 4 to 6 per cent.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a sulphite (of sodium, for instance—made by passing sulphurous acid into solution of carbonate of sodium) add a drop or two of dilute hydrochloric acid; sulphurous acid gas escapes, recognized by a peculiar pungent smell.

This smell is the same as that evolved on burning lucifer matches that have been tipped with sulphur. It is due, probably, not to the gas ($SO_2$), but to sulphurous acid ($H_2SO_3$) formed by the union of sulphurous acid gas with either the moisture of the air or that on the surface of the mucous membrane of the nose. It is highly suffocating.

*Second Analytical Reaction.* — To a sulphite add a little water, a fragment or two of zinc, and then hydrochloric acid; sulphuretted hydrogen will be evolved, known by its putrid odor and action on a piece of paper placed like a cap on the mouth of the test-tube, and moistened with a drop of solution of acetate of lead, black sulphide of lead being formed. Sulphurous acid may be detected in acetic acid, or in hydrochloric acid, by this test.

$$H_2SO_3 + H_6 = H_2S + 3H_2O.$$

### *Other Aalytical Reactions.*

To solutions of neutral sulphites add nitrate or chloride of barium, chloride of calcium, or nitrate of silver; in each case white sulphites of the various metals are precipitated. The barium sulphite is soluble in weak hydrochloric acid; but if a drop or two of chlorine-water is first added, barium sulphate is formed, which is insoluble in acids. The other precipitates are also soluble in acids. The silver sulphite is decomposed on boiling, sulphuric acid being formed, and metallic silver set free.

To recognize the three radicals in an aqueous solution of sulphides, sulphites, and sulphates, add chloride of barium, filter, and wash the precipitate. In the filtrate sulphides are detected by the sulphuretted hydrogen evolved on adding an acid. In the precipitate sulphites are detected by the odor of sulphurous acid produced on adding hydrochloric acid, and sulphates by their insolubility in the acid.

## QUESTIONS AND EXERCISES.

508. What are the differences between sulphurous acid and sulphurous acid gas, sulphites and acid sulphites?

509. State the characters of sulphurous acid gas.

510. How is the official Sulphurous Acid prepared?

511. By what test may sulphurous acid be recognized in acetic acid?

512. Give a method by which sulphites may be detected in presence of sulphides and sulphates.

---

## SULPHURIC ACID AND OTHER SULPHATES.

Formula of Sulphuric Acid $H_2SO_4$. Molecular weight 98.

Sulphates occur in nature; but the common and highly important hydrogen sulphate, sulphuric acid, is made artificially.

*Preparation of the acid. General nature of the Process.*—Sulphur itself, or sometimes the sulphur in iron pyrites, is first converted into sulphurous acid gas by burning in air, and this gas, by moisture and oxygen, into sulphuric acid ($SO_2 + H_2O + O = H_2SO_4$).

*Details of the process.*—The oxygen necessary to oxidize the sulphurous acid gas cannot directly be obtained from air, but indirectly, the agency of nitric oxide ($NO$) being employed—this gas becoming nitric peroxide ($NO_2$) by the action of the air, and the nitric peroxide again becoming nitric oxide by the action of the sulphurous acid gas, and so on. A small quantity of nitric oxide gas will in this way act as carrier of oxygen from the air to very large quantities of sulphurous acid.

The following equations represent the successive steps:—

$$\underset{\text{Sulphur.}}{S_2} + \underset{\text{Oxygen (of the air).}}{2O_2} = \underset{\text{Sulphurous acid gas.}}{2SO_2}$$

$$\underset{\text{Sulphurous acid gas.}}{SO_2} + \underset{\text{Water.}}{H_2O} = \underset{\text{Sulphurous acid.}}{H_2SO_3}$$

$$\underset{\text{Nitric oxide.}}{2NO} + \underset{\text{Oxygen (of the air).}}{O_2} = \underset{\text{Nitric peroxide.}}{2NO_2}$$

---

$$\underset{\text{Sulphurous acid.}}{H_2SO_3} + \underset{\text{Nitric peroxide.}}{NO_2} = \underset{\text{Sulphuric acid.}}{H_2SO_4} + \underset{\text{Nitric oxide.}}{NO}$$

On the large scale the sulphurous acid gas is produced by burning sulphur in furnaces; it is carried, together with the nitric vapors, by flues into leaden chambers, where jets of steam supply the necessary moisture; the steam also, condensing, prevents other reactions.

The resulting dilute sulphuric acid is concentrated by evaporation in leaden vessels.

The nitric peroxide is in the first instance obtained from nitric acid, and this from nitrate of potassium or sodium by the action of a small quantity of the sulphuric acid of a previous operation.

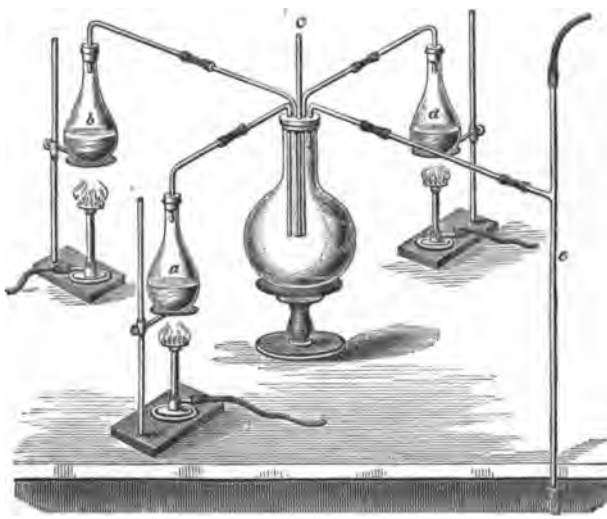
$$\underset{\text{Nitrate of sodium.}}{2NaNO_3} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Sulphate of sodium.}}{Na_2SO_4} + \underset{\text{Nitric acid.}}{2HNO_3}$$

$$\underset{\text{Sulphurous acid.}}{3H_2SO_3} + \underset{\text{Nitric acid.}}{2HNO_3} = \underset{\text{Sulphuric acid.}}{3H_2SO_4} + \underset{\text{Water.}}{H_2O} + \underset{\text{Nitric oxide.}}{2NO}$$

*Other processes.*—Sulphuric acid may be obtained by other processes, as by distilling the sulphate of iron resulting from the natural oxidation of iron pyrites by air; but it is seldom so made at the present day. The sulphate of iron was formerly called *green vitriol* (p. 143), and the distilled product *oil of vitriol;* the latter in allusion to its consistence and origin.

*Experiment.*—For purposes of practical study, a small quantity may be made (as shown in fig. 39) by passing, *a*, sulphurous acid gas

Fig. 39.




Experimental Manufacture of Sulphuric Acid.

(p. 306), *b*, nitric oxide in small quantity (p. 289), *c*, air (forced through by aid of bellows or a gas-holder or drawn through by a *water-aspirator*, *e*), and occasionally, *d*, steam (generated in a

Florence flask) through glass tubes, nearly to the bottom of a two- or three-quart flask.

$$SO_2 + H_2O = H_2SO_3; \quad | \quad 2NO + O_2 = 2NO_2;$$
$$H_2SO_3 + NO_2 = H_2SO_4 + NO.$$

A slow current of sulphurous acid gas, air, and steam, and a small quantity of nitric oxide, will furnish, in the course of a few minutes, enough sulphuric acid for recognition by the first of the following analytical reactions.

*Purification.*—Sulphuric acid may contain arsenic, nitrous compounds, and salts. Arsenic may be detected by the hydrogen-test (p. 173), nitrous compounds by powdered sulphate of iron (which acquires a violet tint if they are present), and salts by the residue left on boiling a little to dryness in a crucible in a fume-chamber. If only nitrous compounds are present, the acid may be purified by heating with about one-half per cent. of sulphate of ammonium—water and nitrogen being produced (Pelouze.) If arsenic occurs, boil with a small quantity of hydrochloric acid, which converts the arsenic into chloride of arsenicum; or heat with a little nitric acid, which converts arsenious ($As_2O_3$) into arsenic anhydride ($As_2O_5$), then add sulphate of ammonium, and distil in a retort containing pieces of quartz and heat by an annular-shaped burner (to prevent "bumping," see p. 281). The arsenic anhydride remains in the retort. (Arsenious anhydride would be carried over with the sulphuric acid vapors.) By distillation the acid is also purified from salts (such as $NaHSO_4$) which are not volatile.

*Quantivalence.*—The sulphuric radical being bivalent ($SO_4''$), acid as well as normal sulphates may exist. Acid sulphate of potassium ($KHSO_4$) is an illustration of the former, sulphate of sodium ($Na_2SO_4$) of the latter; *double sulphates* may also occur, such as that of potassium and magnesium ($K_2SO_4,MgSO_4,6H_2O$). Sulphates generally contain water of crystallization.

*Pure sulphuric acid* ($H_2SO_4$) is of specific gravity 1.848. The best "oil of vitriol" of commerce, a colorless liquid of oily consistence, is of specific gravity 1.843, and contains 96.8 per cent. of real acid ($H_2SO_4$). This is the *Acidum Sulphuricum*, B. P. and U. S. P. The *Acidum Sulphuricum Dilutum*, B. P., sp. gr. 1.094 (U. S. P. 1.082) contains about 13½ (13.64 per cent. of acid ($H_2SO_4$); and the *Acidum Sulphuricum Aromaticum*, B. P. and U. S. P., a dilute acid in which are dissolved the soluble aromatic parts of cinnamon and ginger, also contains nearly 13½ (11.36) per cent. of acid ($H_2SO_4$). There are some definite compounds of sulphuric acid with water; the first ($H_2SO_4,H_2O$) may be obtained in crystals.

*Sulphuric anhydride* ($SO_3$) is a white silky crystalline solid, having no acid properties. It is made by distilling sulphuric acid with phosphoric anhydride ($3H_2SO_4 + P_2O_5 = 2H_3PO_4 + 3SO_3$). It appears to unite with sulphuric acid and some other normal sulphates to form compounds ($R'_2SO_4,SO_3$) resembling in constitution red chromate of potassium or borax. The fuming sulphuric acid ($H_2SO_4,SO_3$), made at Nordhausen in Saxony, seems to be such a body.

*Note.*—Sulphuric acid is a most valuable compound to all chemists and manufacturers of chemical substances. By its agency, direct or indirect, many, if not most, chemical transformations are effected. To describe all its uses would be to write a work on chemistry.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To solution of a sulphate add solution of a barium salt; a white precipitate of sulphate of barium ($BaSO_4$) falls. Add nitric acid and boil the mixture; the precipitate does not dissolve.

This reaction is as highly characteristic of sulphate as it has been stated to be of barium salts (*vide* page 103). The only error likely to be made in its application is that of overlooking the fact that nitrate and chloride of barium are less soluble in strong acid than in water. On adding the barium salt to the acid liquid, therefore, a white precipitate may be obtained, which is simply the nitrate or chloride of barium. The appearance of such a precipitate differs considerably from that of the barium sulphate; hence a careful operator will not be misled. Should any doubt remain, water should be added, which will dissolve the nitrate or chloride, but not affect the sulphate.

*Second Analytical Reaction.*—Mix a fragment of an insoluble sulphate ($BaSO_4$ *e.g.*) with carbonate of potassium or of sodium; or, better, with both carbonates, and fuse the mixture in a small crucible. Digest the residue when cold, in water, and filter; the filtrate may be tested for the sulphuric radical.

This is a convenient method of qualitatively analyzing insoluble sulphates, such as those of barium and lead.

*Third Analytical Reaction.*—Mix a fragment of an insoluble sulphate with a little alkaline carbonate on a piece of charcoal, taking care that some of the charcoal-dust is included in the mixture. Heat the little heap in the blowpipe-flame until it fuses, and, when cold, add a drop of acid; sulphuretted hydrogen is evolved, recognized by its odor.

This is another process for the recognition of insoluble sulphates. Other preparations of sulphur, and sulphur itself, give a similar result. It is therefore rather a test for sulphur and its compounds than sulphates only; but the absence of other salts can generally, if necessary, be previously determined.

*Note.*—The presence of the sulphuric radical in a solution having been proved by the above reactions, its occurrence as the normal sulphate of a metal is demonstrated by the neutral, or nearly neutral, deportment of the liquid with test-paper, and the detection of the metal—its occurrence as sulphuric acid or an acid sulphate by the sourness of the liquid to the taste, and the effervescence produced on the addition of a carbonate.

*Antidote.*—In cases of poisoning by strong sulphuric acid, solution of carbonate of sodium (common washing-soda), magnesia and water, etc., may be administered as antidotes.

## QUESTIONS AND EXERCISES.

513. What is the formula of sulphuric acid, and what its molecular weight?

514. How is it related to other sulphates?

515. Write a short article on the manufacture of sulphuric acid, giving either diagrams or equations.

516. How may nitrous compounds be detected in, and eliminated from, sulphuric acid?

517. State the method by which the presence of arsenic is detected in sulphuric acid, and explain the process by which it may be removed.

518. Define sulphates, acid sulphates, and double sulphates.

519. What percentage of real acid is contained in commercial oil or vitriol?

520. State the strength of the official "diluted" and "aromatic" sulphuric acid.

521. By what process is sulphuric anhydride obtained from Nordhausen sulphuric acid?

522. Explain the reactions which occur in testing for sulphates.

523. Ascertain by calculation the weight of oil of vitriol (of 96.8 per cent.) necessary for the production of one ton of dry sulphate of ammonium.—*Ans.* 1718 pounds.

524. Name the antidotes in cases of poisoning by strong sulphuric acid.

## CARBONIC ACID AND OTHER CARBONATES.

Formula of carbonic acid $H_2CO_3$. Molecular weight 62. Formula of carbonic acid gas, or carbonic anhydride, commonly termed carbonic acid, $CO_2$.

*Sources.*—Carbonates (compounds containing the grouping $CO_3$) are very common in nature, the calcium carbonate ($CaCO_3$) being widely distributed as chalk, limestone, or marble. The hydrogen carbonate, true carbonic acid, is not known, unless indeed carbonic acid gas assumes that condition on dissolving in water (*Aqua Acidi Carbonici*, U. S. P.). Such a solution (see page 85) changes the color of blue litmus paper, and the gas does not; this may be because only the true acid ($H_2CO_3$) affects the litmus, or because the gas ($CO_2$) cannot come into real contact with the litmus without a medium. From the commonest natural carbonate, carbonate of calcium, are derived the carbonic constituents of the one most frequently used in medicine, carbonate of sodium.

*Carbonate of sodium* is prepared from the chief natural salt, the chloride. After the chloride has been converted into sulphate (salt-cake) by sulphuric acid (or by sulphurous acid, air, and steam—Hargreaves modification)—

$$2NaCl + H_2SO_4 = Na_2SO_4 + 2HCl,$$

the sulphate is roasted with limestone and small coal, by which carbonate of sodium and sulphide of calcium are formed:

$$Na_2SO_4 + C_4 + CaCO_3 = CaS + Na_2CO_3 + 4CO.$$

Carbonic oxide gas and a little carbonic acid gas from the excess of chalk escape; the residual mass (black ash) is digested in water, in which the carbonate of sodium dissolves, the sulphide of calcium with a little oxide remaining insoluble. The solution is evaporated to dryness, and yields crude carbonate of sodium. This is roasted with a small quantity of sawdust, to convert any caustic soda, resulting from the action of the lime on the carbonate, into normal carbonate. The product is *soda-ash*. Dissolved in water and crystallized, it constitutes the ordinary "soda" used for washing purposes; recrystallized and sometimes ground, it forms the official carbonate of sodium (*Sodii Carbonas*, U. S. P.) ($Na_2CO_3,10H_2O$). The reaction is rendered more intelligible by regarding it as occurring in two stages: 1st, the reduction of the sulphate of sodium to sulphide by the carbon of the coal—

$$Na_2SO_4 + C_4 = Na_2S + 4CO;$$

2d, the reaction of the sulphide of sodium and carbonate of calcium, giving soluble carbonate of sodium, thus—

$$Na_2S + CaCO_3 = Na_2CO_3 + CaS.$$

The sulphur in the resiual sulphide of calcium may be recovered by exposure to air, and the subsequent action of hydrochloric acid. Some hyposulphite of calcium ($CaS_2O_3$) is first formed, and the action of the acid on this and on undecomposed sulphide gives chloride of calcium, water, and sulphur.

*Another process.* — To a strong solution of common salt, bicarbonate of ammonium is added, when a precipitate of bicarbonate of sodium occurs. The resulting chloride of ammonium may be converted into carbonate by heating with chalk, and the carbonate be more fully carbonated by carbonic acid gas obtained by heating the bicarbonate of sodium, which is thereby reduced to the ordinary neutral carbonate.

*Carbonic acid gas* ($CO_2$) is a product of the combustion of all carbonaceous matters. It is constantly exhaled by animals and inhaled by plants, its intermediate storehouse being the atmosphere, throughout which it is equally distributed by *diffusion* (*vide* p. 24) to the extent of about 4 parts in 10,000. A larger proportion than that just mentioned gives to confined air depressing effects, 4 or 5 per cent. rendering the atmosphere poisonous when taken into the blood from the lungs. Carbonic acid, however, may be taken into the stomach with beneficial sedative effects; hence, probably, much

of the value of such effervescing liquids as soda-water, lemonade, and solutions of the various granulated preparations and effervescing powders (*vide* p. 87). The gas liquefies on being compressed, and the liquid solidifies on being evolved. Carbonic acid gas is twenty-two times as heavy as hydrogen, and about half as heavy again as air.

## Reactions.

*Synthetical and Analytical Reactions.*—1. To a fragment of marble in a test-tube add water and then hydrochloric acid; carbonic acid gas ($CO_2$) is evolved, and may be conveyed into water or solutions of salts by the usual delivery-tube.

This is the process of the British Pharmacopœia, and the one usually adopted for experimental purposes. Passed into carbonate of sodium, the gas gives *Sodii Bicarbonas* (p. 83), and into carbonate of potassium, *Potassii Bicarbonas* (p. 71). On the large scale the gas is prepared from chalk or marble and sulphuric acid, frequent stirring promoting its escape.

2. Pass the gas into lime-water; a white precipitate of carbonate of calcium ($CaCO_3$) falls. Solution of subacetate of lead may be used instead of, and is perhaps even a more delicate test than, lime-water.

The evolution of a gas on adding an acid to a salt, warming the mixture if necessary, the gas being inodorous and giving a white precipitate with lime-water, is sufficient evidence of the presence of a carbonate. Carbonates in solution of ammonia, potash, or soda, may be detected by the direct addition of solution of lime.

3. Blow air from the lungs through a glass tube into lime-water; the presence of carbonic acid gas is at once indicated.

The passage of a considerable quantity of normal air through lime-water produces a similar effect. A bottle containing lime-water soon becomes coated with carbonate of calcium from absorption of atmospheric carbonic acid gas.

4. Fill a dry test-tube with the gas, by pressing the delivery-tube of the above apparatus to the bottom of the test-tube. Being rather more than once and a half as heavy as the air (1.529) it will displace the latter. Prove the presence of the gas by pouring it slowly, as if a visible liquid, into another test-tube containing lime-water; the characteristic cloudiness and precipitate are obtained on gently shaking the lime-water.

In testing for carbonates by bringing evolved gas into contact with lime-water, the preparation and adaptation of a delivery-tube may often be avoided by pouring the gas from the generating-tube into that containing the lime-water in the manner just indicated.

5. Pass carbonic acid gas through lime-water until the precipitate at first formed is dissolved. The resulting liquid is a solution of carbonate of calcium in carbonic acid water. Boil the solution; carbonic acid gas escapes, and the carbonate is again precipitated.

This experiment will serve to show how chalk is kept in solution in ordinary well-waters, giving the property of "hardness," and how the *fur* or stone-like deposit in tea-kettles and boilers is formed. It should be here stated that sulphate of calcium produces similar hardness, and that these, with small quantities of the sulphate and carbonate of magnesium, constitute the hardening constituents of well-waters, a curd (oleate of calcium or magnesium) being formed whenever soap is used with such waters. An enormous amount of soap is wasted through the employment of hard water for washing-purposes. The hardness produced by the earthy carbonates is termed "temporary hardness," because removable by ebullition; that by the earthy sulphates "permanent hardness" because unaffected by ebullition. The addition of lime-water or a mixture of lime and water removes temporary hardness (reac. 2, page 314), and carbonate of sodium, "washing soda," both temporary and permanent hardness, in the latter case sulphate of sodium remaining in solution. Carbonate of barium (ground witherite) also decomposes sulphates of calcium and magnesium, sulphate of barium being precipitated and carbonates of calcium or magnesium formed; the latter and the carbonates originally in the water may then be precipitated by ebullition or by the action of lime-water. But the injurious effects of barium salts on man and the lower animals prevent the carbonate being used for purifying water for drinking purposes, as by accident or an unforeseen reaction a portion might become dissolved.

---

## QUESTIONS AND EXERCISES.

525. Name the chief natural carbonates.

526. What are the formulæ of carbonic acid and carbonic acid gas?

527. Adduce evidence of the existence of true carbonic acid.

528. Trace the steps by which the carbonic constituent of chalk is transferred to sodium by the process usually adopted in alkali-works —the manufacture of "soda."

529. Carbonic acid gas is constantly exhaled from the lungs of animals; why does it not accumulate in the atmosphere?

530. What is the effect of pressure on carbonic acid gas?

531. State the specific gravity of carbonic acid gas.

532. By what processes may carbonic acid gas be obtained for experimental and manufacturing-purposes?

533. Describe the action of carbonic acid gas on the carbonates of potassium or sodium.

534. How may carbonic acid be detected in expired air?

535. To what extent is carbonic acid gas heavier than air?

536. What quantity of chalk (90 per cent. pure) will be required to furnish the carbonic acid necessary to convert one ton of carbonate of potassium (containing 83 per cent. of $K_2CO_3$) into acid carbonate, supposing no gas to be wasted?—*Ans.* 1500 lbs.

537. Define "hardness" in water.

538. How may the presence of carbonates be demonstrated?

---

## OXALIC ACID AND OTHER OXALATES.

Formula of Oxalic Acid $H_2C_2O_4,2H_2O$. Molecular weight 126.

*Source* —Oxalates occur in nature in the juices of some plants, as wood-sorrel, rhubarb, the common dock, and certain lichens; but the hydrogen oxalate (oxalic acid) and other oxalates are all made artificially. The carbon of many organic substances yields oxalic acid when those substances are boiled with nitric acid, and an alkaline oxalate when they are roasted with a mixture of the hydrates of potassium and sodium.

*Experimental Process.*—On the small scale, a mixture of nitric acid, 10 parts, loaf sugar 2 parts, and water 3 parts, quickly yields the acid. Abundance of red fumes are at first evolved. On cooling, crystals are deposited. A more dilute acid, kept warm, acts more slowly but yields a larger product.

*Manufacturing Process.*—On the large scale, sawdust is roasted with alkalies, resulting oxalate of sodium decomposed by lime with formation of oxalate of calcium, the latter digested with sulphuric acid, and the liberated oxalic acid (Oxalic Acid of Commerce, B. P.) made commercially pure by recrystallization (Oxalic Acid, Purified, B. P., *Acidum Oxalicum*, U. S. P.).

*Chemically pure oxalic acid.*—The acid made from sugar, recrystallized two or three times, is quite pure. Commercial acid should be mixed with insufficient water for complete solution, and the mixture occasionally shaken. Impurities remain undissolved, and the saturated solution evaporated yields crystals which seldom require to be recrystallized.

*Quantivalence.*—The elements represented by the formula $C_2O_4$ are those characteristic of oxalates. They form a bivalent grouping; hence normal oxalates ($R'_2C_2O_4$) and acid oxalates ($R'HC_2O_4$) exist.

*Salt of sorrel* is a crystalline compound of oxalic acid with acid potassium oxalate, the crystals containing two molecules of water of crystallization ($KHC_2O_4,H_2C_2O_4,2H_2O$).

*Oxalate of iron* (*Ferri Oxalas*, U. S. P.) is a crystalline powder made by precipitating a solution of sulphate of iron with oxalic acid. When heated in contact with air it decomposes with a faint combustion, and leaves a residue of not less than forty-eight per cent. of red oxide of iron.

*Analytical Reactions (Tests).*

*First Analytical Reaction.*—To solution of an oxalate (oxalate of ammonium, *e. g.*) add solution of chloride of calcium; a white precipitate falls. Add to the precipitate excess of acetic acid; it is insoluble. Add hydrochloric acid; the precipitate is dissolved.

The formation of a white precipitate on adding a calcium or barium salt, insoluble in acetic but soluble in hydrochloric or nitric acid, is usually sufficient proof of the presence of an oxalate. The action of the liquid on litmus paper, effervescence with carbonate of sodium, and absence of metals, would indicate that the oxalate is that of hydrogen, oxalic acid.

*Note.*—The *barium* oxalate *is* slightly soluble in acetic acid (Souchay and Lenssen), and enough may be dissolved by this acid from a mixed barium precipitate (produced on adding chloride or nitrate of barium to a solution of mixed salts) to give the above reaction on adding chloride of calcium to the filtered acetic liquid—an effect sometimes useful in the analysis of mixed substances (Davies).

*Antidote.*—In cases of poisoning by oxalic acid or salt of sorrel, chalk and water may be administered as a chemical antidote (with the view of producing the insoluble oxalate of calcium), emetics and the stomach-pump being used as soon as possible.

*Second Analytical Reaction.*—Heat a fragment of any common fixed metallic oxalate (an oxalate of potassium for example) in a test-tube; decomposition occurs, carbonic oxide (CO) (a gas that will be noticed subsequently) is liberated, and a carbonate of the metal remains. Add water and then an acid to the residue; effervescence occurs.

This is a ready test for ordinary insoluble oxalates, and is trustworthy if, on heating the substance, no charring occurs, or not more than gives a gray color to the residue. Organic salts of metals decompose when heated, and leave a residue of carbonate, but except in the case of oxalate, the residue is always accompanied by much charcoal. Insoluble oxalates and organic salts of such metal as lead and silver are, of course, liable to be reduced to oxide or even metal by heat. Such oxalates may be decomposed by boiling with solution of carbonate of sodium, filtering, and testing the filtrate for oxalates by the chloride of calcium test.

*Other Analytical Reactions.*—Nitrate of silver gives, with oxalates, white oxalate of silver ($Ag_2C_2O_4$).——Dry oxalates are decomposed when heated with strong sulphuric acid, carbonic oxide and carbonic acid gases escaping. If much of the substance be operated on, the gas may be washed with an alkali, the carbonic acid be thus removed, and the carbonic oxide be ignited; it will be found to burn with a characteristic bluish

flame.——Oxalates, when mixed with water, black oxide of manganese (free from carbonates), and sulphuric acid, yield carbonic acid gas, which may be tested by lime-water in the usual manner.——Not only such insoluble oxalates as those of lead and silver, above referred to, but any common insoluble oxalates, such as those of calcium and magnesium, may be decomposed by ebullition with solution of carbonate of sodium; after filtration the oxalic radical will be found in the clear liquid as soluble oxalate of sodium.

*Test of Purity.*—"Purified oxalic acid . . . . is entirely dissipated by a heat below 350° F." (B. P.)

## QUESTIONS AND EXERCISES.

539. Explain the constitution of oxalates.
540. State how oxalates are obtained.
541. What is the quantivalence of the oxalic radical?
542. Give the formula of "salt of sorrel."
543. Mention the chief test for oxalic acid and other soluble oxalates.
544. Name the antidote for oxalic acid, and describe its action.
545. By what reactions are insoluble oxalates recognized?

## TARTARIC ACID AND OTHER TARTRATES.

Formula of Tartaric Acid $H_2C_4H_4O_6$, or $H_2\overline{T}$.
Molecular weight 150.

*Source.*—Tartrates exist in the juice of many fruits; but it is from that of the grape that our supplies are usually obtained. Grape-juice contains much acid tartrate of potassium, which is gradually deposited when the juice is fermented, as in making wine; for acid tartrate of potassium, not very soluble in aqueous liquids, is still less so in spirituous, and hence crystallizes out as the sugar of the grape-juice is gradually converted into alcohol. It is found with tartrate of calcium lining the vessels in which wine is kept; and it is from this crude substance, termed *argal* or *argol*, also from the albumenoid yeasty matter or "lees" deposited at the same time, as well as from what tartrate may be remaining in the marc left after the juice has been pressed from the grapes, that by rough crystallization "tartar" is obtained. From the latter tartaric acid and other tartrates are prepared. In old dried grapes (Raisins; *uvæ*, B. P.) crystalline masses of tartar and of grape sugar are constantly met with.

*Cream of Tartar*, purified by crystallization (*Potassæ Tartras Acida*, B. P., *Potassii Bitartras*, U. S. P.), occurs as a "gritty white powder, or fragments of cakes crystallized on one surface;" of

a pleasant acid taste, soluble in 180 parts of cold and 6 of boiling water, insoluble in spirit.*

*Quantivalence.*—The elements represented by the formula $C_4H_4O_6$ are those characteristic of tartrates. They form a bivalent grouping; hence normal tartrates ($R'_2\overline{T}$) and acid tartrates ($R'H\overline{T}$) exist. Tartrate of potassium, the *Potassii Tartras* of the U. S. Pharmacopœia ($K_2C_4H_4O_6$), and Rochelle Salt, or tartrate of potassium and sodium ($KNaC_4H_4O_6,4H_2O$), the official *Potassii et Sodii Tartras* (*Soda Tartarata*, B. P.), are illustrations of normal tartrates, while Cream of Tartar is an example of acid tartrates. The only official tartrate not apparently included in these general formulæ is tartar-emetic (*Antimonium Tartaratum*, B. P., *Antimonii et Potassii Tartras*, U. S. P.), which is sometimes regarded as the double tartrate of potassium and a hypothetical radical, antimonyl (SbO), thus, $KSbOC_4H_4O_6$. Probably, however, it is but an oxytartrate of antimony ($Sb_2O_2\overline{T}$) with normal tartrate of potassium ($K_2\overline{T}$); for there are several oxycompounds of antimony analogous to the oxycompounds of bismuth that have been described (p. 252), normal salts partially decomposed by water into oxides, and many of these oxycompounds readily unite with normal salts of other basylous radicals. Tartar emetic would thus be oxytartrate of antimony, with tartrate of potassium ($Sb_2O_2\overline{T},K_2\overline{T}$, or $Sb_2O_2C_4H_4O_6,K_2C_4H_4O_6$.

## Tartaric Acid.

*Tartaric Acid* (*Acidum Tartaricum*, B. P. and U. S. P.) is obtained, according to the British Pharmacopœia, by boiling cream of tartar (*Potassæ Tartras Acida*, B. P., *Potassii Bitartras*, U. S. P.) with water, adding chalk till effervescence ceases, and then chloride of calcium so long as a precipitate falls; the two portions of tartrate of calcium thus consecutively formed are thoroughly washed, treated with sulphuric acid, the mixture boiled for a short time, resulting sulphate of calcium mostly separated by filtration, the filtrate concentrated by evaporation, any sulphate of calcium that may have deposited removed as before, and concentration continued until the solution is strong enough to crystallize. Tartrate of calcium from 9 ounces of cream of tartar requires 5 ounces by weight of sulphuric acid for complete decomposition.

---

* A boiling solution of tartar yields a floating crust of minute crystals on cooling—just as milk yields a floating layer of cream, hence the term *cream* of tartar.

"It is called *tartar*," says Paracelsus, "because it produces oil, water, tincture, and salt, which burn the patient as tartarus does." *Tartarus* is Latin (Τάρταρος, Tartaros, Greek) for *hell*. The products of its destructive distillation are certainly somewhat irritating in taste and smell; and the "salt" (carbonate of potassium) that is left is diuretic, and, in larger quantities, powerfully corrosive.

$$\underset{\text{Acid tartrate of potassium.}}{2KH\overline{T}} + \underset{\text{Carbonate of calcium.}}{CaCO_3} = \underset{\text{Tartrate of calcium.}}{Ca\overline{T}} + \underset{\text{Tartrate of potassium.}}{K_2\overline{T}} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

$$\underset{\text{Tartrate of potassium.}}{K_2\overline{T}} + \underset{\text{Chloride of calcium.}}{CaCl_2} = \underset{\text{Tartrate of calcium.}}{Ca\overline{T}} + \underset{\text{Chloride of potassium.}}{2KCl}$$

$$\underset{\text{Tartrate of calcium.}}{2Ca\overline{T}} + \underset{\text{Sulphuric acid.}}{2H_2SO_4} = \underset{\text{Sulphate of calcium.}}{2CaSO_4} + \underset{\text{Tartaric acid.}}{2H_2\overline{T}}$$

Tartaric acid occurs in colorless crystals, or the same powdered. It is strongly acid and readily soluble in water or spirit. One part in 8 of water and 2 of spirit of wine forms "Solution of Tartaric Acid," B. P. Its aqueous solution is not stable.

Parcels of tartaric acid often contain crystals of an allotropic or physically isomeric modification (*vide* "Allotropy" and "Isomerism" in Index). It is termed *Paratartaric acid* (*παρά*, *para*, beside) or *Racemic acid* (*racemus*, a bunch of grapes), and is a combination of ordinary tartaric acid, whose solution twists a ray of polarized light to the right hand (dextrotartaric or dextroracemic acid), and of lævotartaric or lævoracemic acid, whose solution twists a polarized ray to the left. Racemic acid is inactive in this respect, the opposite properties of its constituents neutralizing each other. Racemic acid is less soluble in alcohol than tartaric acid.

### Reactions.

### Tartrate of Potassium.

*Synthetical Reactions.*—To a small quantity of a strong solution of carbonate of potassium add acid tartrate of potassium so long as effervescence occurs; the resulting liquid is solution of normal tartrate of potassium (*Potassii Tartras*, U. S. P.) ($K_2\overline{T}$), crystals of which may be obtained on evaporation.

*Note.*—This is a common method of converting an acid salt of a bivalent acidulous radical into a normal salt. The carbonate added need not be a carbonate of the same, but may be of a different metal; compounds like Rochelle salt ($KNa\overline{T}$) are then obtained. Thus:—

### Tartrate of Potassium and Sodium.

To a strong hot solution of carbonate of sodium add acid tartrate of potassium until effervescence ceases; the resulting liquid is solution of tartrate of potassium and sodium; on cooling, it yields crystals. This is the official process (*Soda Tartarata*, B. P., *Potassii et Sodii Tartras*, U. S. P.) ($KNaC_4H_4O_6,4H_2O$).

$$\underset{\text{Carbonate of sodium.}}{Na_2CO_3} + \underset{\text{Acid tartrate of potassium.}}{2KHC_4H_4O_6} = \underset{\text{Tartrate of potassium and sodium.}}{2KNaC_4H_4O_6} + \underset{\text{Water.}}{H_2O} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

Crystals of Rochelle salt are usually halves of colorless, transparent, right rhombic prisms, slightly efflorescent in dry air, soluble in five parts of boiling water. Tartrate of potassium is slightly deliquescent, soluble in about four parts of boiling water.

**Equivalent Weights** *of Tartaric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium, Bicarbonate of Sodium, and Carbonates of Ammonium and Magnesium; repeated for* 20 *parts of each* (and, incidentally, for other proportions).

| | | | | | | | | | |
|---|---|---|---|---|---|---|---|---|---|
| Tart. Acid ..... | $H_2C_4H_4O_6$ ..... = 150 | 20 | 18½ | 15 | 10½ | 17½ | 25½ | 31½ |
| Carb. Potas. ..... | $K_2CO_3$ (of 84 per cent.) = 164 | 22 | 20 | 16½ | 11½ | 19½ | 28 | 34½ |
| Bicarb. Pot. ..... | $2(KHCO_3)$ ..... = 200 | 26⅔ | 24⅔ | 20 | 14 | 3⅓ | 34 | 42 |
| Carb. Soda (cryst.) | $Na_2CO_3,10H_2O$ ..... = 286 | 38 | 34⅓ | 28½ | 20 | 34 | 48½ | 60 |
| Bicarb. Soda..... | $2(NaHCO_3)$. ..... = 168 | 22½ | 20 | 16⅔ | 11⅔ | 20 | 28½ | 35⅓ |
| Carb. Ammon. ..... | $(N_4H_{16}C_2O_6)+2$..... = 118 | 15¾ | 14⅓ | 11⅔ | 8¼ | 14 | 20 | 24⅔ |
| Carb. Magnes. ..... | $(MgCO_3)_3Mg2HO,4H_2O$ =95.5 | 12¾ | 11½ | 9⅔ | 6⅔ | 11⅓ | 16 | 20 |

Thus 20 parts (grains or other weights) of tartaric acid neutralize 22 of carbonate of potassium, 26⅔ of bicarbonate of potassium, 38 of carbonate of sodium, 22½ of bicarbonate of sodium, 15¾ of carbonate of ammonium, or 12¾ of carbonate of magnesium. Other quantities of tartaric acid (18½, 15, 10½, 16⅔, 25½, 31½) saturate the amounts of salts memtioned in the other columns and *vice versâ*. A similar Table for Citric Acid will be found on page 325, and for both acids in the Appendix. These Tables afford good illustrations of the laws of chemical combination (page 47). The reader should verify a few of the numbers by calculation from the atomic weights of the elements concerned in the reactions, remembering that the salts formed are considered to be neutral in constitution. In medical practice effervescing saline draughts are often designedly prescribed to contain an amount of acid or alkali considerably in excess of the proportions required for perfect neutrality.

A common form of *Seidlitz Powder* consists of 3 parts of Rochelle salt (120 grains) with 1 (40 grains) of acid carbonate of sodium (the mixture usually wrapped in blue paper), and 1 (40 grains) of tartaric acid (wrapped in white paper). When administered, the latter is dissolved in a tumbler rather more than half full of water, the former added, and the mixture drunk during effervescence. It will be seen that the salts swallowed are tartrate of potassium and sodium ($KNa\overline{T},4H_2O$), tartrate of sodium ($Na_2\overline{T},2H_2O$), and acid tartrate of sodium or potassium. The last-mentioned salt results because 11½ per cent. (4½ grains) of the tartaric acid is in excess of the quantity

necessary for the formation of neutral tartrate of sodium. This amount of acid salt gives agreeable acidity to the draught. The United States formula (*Pulveres Effervescentes Aperientes*, U. S. P.) includes rather less tartaric acid, so that only neutral salts are formed.

### *Analytical Reactions* (*Tests*).

*First Analytical Reaction.* — To solution of any normal tartrate, or tartaric acid made neutral by solution of soda, add solution of chloride of calcium; a white precipitate, tartrate of calcium, falls. Collect the precipitate on a filter, wash, place a small quantity in a test-tube, and add solution of potash; on stirring the mixture the precipitate dissolves. Heat the solution; the tartrate of calcium is again precipitated.

In the above reaction a fair amount of the chloride of calcium solution should be added at once, and the whole test performed without delay, or the calcium tartrate will assume a crystalline character, and be with difficulty dissolved by the potash.

The solubility of tartrate of calcium in cold potash solution enables the analyst to distinguish between tartrates and citrates, otherwise a difficult matter. Citrate of calcium is not soluble, or only to a very slight extent, in the alkali. The absence of much ammoniacal salt must be insured, citrate as well as tartrate of calcium being soluble in solutions of salts of ammonium.

*Second Analytical Reaction.*—Acidulate a solution of a tartrate with acetic acid, add acetate of potassium, and well stir the mixture; a crystalline precipitate of acid tartrate of potassium slowly separates.

This reaction is not applicable in testing for very small quantities of tartrates, the acid tartrate of potassium being not altogether insoluble. The precipitate being insoluble in alcohol, however, the addition of spirit of wine renders the test far more delicate.

*Third Analytical Reaction.*—To a neutral solution of a tartrate add solution of nitrate of silver; a white precipitate of tartrate of silver, $Ag_2C_4H_4O_6$, falls. Boil the mixture; it blackens, owing to the reduction of the salt to metallic silver. Or, before boiling, add a drop, or less, of ammonia; a mirror will form on the tube—adhering well to the glass if the tube was thoroughly cleansed. Even an insoluble tartrate, placed in a dry tube with a few fragments of nitrate of silver and a drop, or less, of ammonia added, gives a mirror-like character to each fragment of the silver salt when the tube is gently rotated some inches above a flame.

*Fourth Analytical Reaction.*—To a neutral or alkaline solution of a tartrate add a few drops of solution of permanganate of potassium, and slowly heat the test-tube; the color is discharged, an oxide of manganese being precipitated. Citrates only reduce the permanganate to green manganate.

*Other Reactions.*—Tartrates heated with strong sulphuric acid char immediately.——Tartaric acid and the soluble tartrates prevent the precipitation of ferric and other hydrates by alkalies, soluble double tartrates being formed (which on evaporation yield liquids that do not crystallize, but, spread on sheets of glass, dry up to thin transparent plates or scales). The *Ferri et Potassii Tartras*, U. S. P. (*Ferrum Tartaratum*, B. P.), is a preparation of this kind.——Tartrates decompose when heated, carbonates being formed and carbon set free, the gaseous products having a peculiar, more or less characteristic smell, resembling that of burnt sugar.

## QUESTIONS AND EXERCISES.

546. State the origin of tartaric acid and other tartrates, and explain the deposition of argol, crude acid tartrate of potassium, during the manufacture of wine.

547. What are the chemical formula and characters of "cream of tartar"?

548. Mention the formula and quantivalence of the tartaric radical.

549. Write formulæ of normal, acid, and double tartrates, tartar-emetic being treated as an oxytartrate of antimony with tartrate of potassium.

550. Give equations or diagrams illustrative of the production of tartaric acid from cream of tartar.

551. By what general process may normal or double tartrates be obtained from acid tartrate of potassium?

552. Work out sums proving the correctness of some of the figures given on p. 321 as showing the saturating power of tartaric acid for various quantities of different carbonates, and give diagrams or equations of the reactions.

553. State the names and quantities of the salts resulting from the admixture of 120 grains of tartrate of potassium and sodium, 40 grains of acid carbonate of sodium, and 40 grains of tartaric acid (Seidlitz powder).

554. Enumerate the tests for tartrates, and explain the effects of heat on tartrates of the metals

## CITRIC ACID AND OTHER CITRATES.

Formula of Citric Acid $H_3C_6H_5O_7,H_2O$ or $H_3\overline{C}iAq$.

Molecular weight 210.

*Source.*—Citric acid (*Acidum Citricum*, B. P. and U. S. P.) exists in the juice of many of our common garden fruits. The pulp of the fruit of *Tamarindus indica* (*Tamarindus*, B. P. and U. S. P.) contains nearly 10 per cent. (in addition to 1.5 of tartaric acid, 5 of malic acid, and 3 per cent. of acid tartrate of potassium). But it is from the lemon or lime that the acid of commerce is usually obtained. For this purpose concentrated lemon-juice is exported from Sicily, concentrated bergamot juice from the Calabrian coast of South Italy, and concentrated lime-juice from the West Indies. The lime-fruit from *citrus bergamia* is official in the Pharmacopœia of India.

*Process.*—The British Pharmacopœia directs that the hot lemon-juice (4 pints) be saturated by powdered chalk, that is, whiting (4½ ounces), the resulting citrate of calcium collected on a filter, washed with hot water till the liquor passes from it colorless (by which not only the coloring matter but the mucilage, sugar, and other constituents of the juice are got rid of), then mixed with cold water (1 pint) decomposed by sulphuric acid (2½ fluidounces in 1½ pint of water), the mixture boiled for half an hour, filtered, the solution evaporated to a density of 1.21, set aside for 24 hours, then poured off from any deposit of crystalline sulphate of calcium, further concentrated, and set aside to crystallize. If the quantity of citrate of calcium to be decomposed is indefinite, the sulphuric acid may be added until a little of the supernatant fluid gives, after a minute or two, a precipitate with solution of chloride of calcium. The concentrated citric solution generally crystallizes very slowly. Shaken violently, however, in a bottle, with a granule or two of solid acid, it quickly yields its citric acid in a pulverulent form, and this, drained and redissolved in a very small quantity of hot water, yields crystals fairly quickly (Warrington).

$$\underset{\text{Citric acid (impure).}}{2H_3C_6H_5O_7} + \underset{\text{Carbonate of calcium.}}{3CaCO_3} = \underset{\text{Citrate of calcium.}}{Ca_3 2C_6H_5O_7} + \underset{\text{Water.}}{3H_2O} + \underset{\text{Carbonic acid gas.}}{3CO_2}$$

$$\underset{\text{Citrate of calcium.}}{Ca_3 2C_6H_5O_7} + \underset{\text{Sulphuric acid.}}{3H_2SO_4} = \underset{\text{Citric acid (pure).}}{2H_3C_6H_5O_7} + \underset{\text{Sulphate of calcium}}{3CaSO_4}$$

*Quantivalence.*—The elements represented by the formula $C_6H_5O_7$ are those characteristic of citrates. They form a trivalent grouping; hence three classes of salts may exist—one, two, or three atoms of the basylous hydrogen in the acid, $H_3C_6H_5O_7$, being displaced by equivalent proportions of other basylous radicals.

*Citric acid* itself is the only citric compound of much direct importance to the pharmacist. It usually occurs in colorless crystals soluble in half their weight of boiling and three-fourths of cold water, less soluble in spirit, and insoluble in ether. A solution of about 34 grains in 1 ounce of water forms a sort of artificial lemon-juice. Citrates heated with strong sulphuric acid to about 212° F. evolve

carbonic oxide gas, and at higher temperatures acetone and carbonic acid gas.

*Action of heat on citric acid.*—Citric acid slowly heated first loses its water of crystallization; afterwards (at 347° F.) the elements of another molecule of water are evolved and a residue obtained from which ether extracts *aconitic acid* ($H_3C_6H_3O_6$), identical with the aconitic acid (and the acid first termed *equisetic*) in various species of *Aconitum* and *Equisetum*.

*The official Lemon Juice* (*Succus Limonum*, B. P., *Limonis Succus*, U. S. P) is to be freshly expressed from the ripe fruit, and contain an average of 32.5 grains of citric acid in 1 fluidounce. The acidity may be ascertained by adding solution of potash or soda (the strength of which has been previously determined with pure crystals of citric acid) till red litmus-paper is fairly turned blue. Before applying this test to commercial specimens, the absence of notable quantities of sulphuric, hydrochloric, acetic, tartaric, or other acid must be insured by application of appropriate reagents. (See also "Lemon Juice," in Index.)

*Mistura Potassii Citratis*, U. S. P., is lemon-juice completely neutralized by bicarbonate of potassium. It is a slightly impure but flavored solution of citrate of potassium.

**Equivalent Weights** *of Citric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium, Bicarbonate of Sodium, and Carbonates of Ammonium and Magnesium; repeated for* 20 *parts of each* (and, incidentally, for other proportions).

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| Citric Acid..... | $H_3C_6H_5O_7,H_2O$ .....= 210 | 20 | 17 | 14 | 9½ | 16½ | 23½ | 29½ |
| Carb. Potas. ..... | ($K_2CO_3$; of 84 p.ct.)+2X3= 246½ | 23½ | 20 | 16½ | 11½ | 19½ | 28 | 34½ |
| Bicarb. Pot. ..... | 3($KHCO_3$) .....= 300 | 28½ | 24½ | 20 | 14 | 24 | 34 | 41½ |
| Carb Sod.(cryst.) | ($Na_2CO_3$10$H_2O$)+2X3 ..= 429 | 40 | 34½ | 28½ | 20 | 34½ | 48½ | 60 |
| Bicarb. Sod. ..... | 3($NaHCO_3$) .....= 252 | 24 | 20½ | 16½ | 11½ | 20 | 28½ | 35 |
| Carb. Ammon. ... | ($N_4H_{16}C_3O_8$)+4X3 .....= 177 | 16¾ | 14¼ | 11¾ | 8¼ | 14 | 20 | 24¾ |
| Carb. Magnes. ... | ($MgCO_3$)$_3$$Mg2HO$,4$H_2O$ +8X3= 143¼ | 13¼ | 11¾ | 9¼ | 6¾ | 11¼ | 16¼ | 20 |

Thus 20 parts (grains, or other weights) of citric acid neutralize 23½ of carbonate of potassium, 28½ of bicarbonate of potassium, 40 of carbonate of sodium, 24 of bicarbonate of sodium, 16¾ of carbonate of ammonium, or 13¼ of carb. of magnesium. Other quantities of acid (17, 14, 9¾, 16¾, 23¾, 29¼) saturate the amount of salts mentioned in the other columns, and *vice versâ*.

This Table, the similar one for tartaric acid (p. 321), and that for both acids (*vide* Appendix) afford good illustrations of some of the laws of chemical combination (p. 47). The reader should verify a few of the numbers by calculation from the atomic weights of the elements concerned in the reactions, remembering that the salts

formed are considered to be neutral in constitution. In medical practice, effervescing saline draughts are often designedly prescribed to contain an amount of acid or alkali considerably in excess of the proportions required for perfect neutrality.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—To a dilute solution of any neutral citrate, or citric acid carefully neutralized by alkali, add solution of chloride of calcium and boil; a white precipitate, citrate of calcium ($Ca_3\overline{Ci}_2$), falls. Treat the precipitate, as for tartrate of calcium (p. 322); it is not perceptibly dissolved by the potash.

A mixture of citrates and tartrates can be separated by this reaction. They are precipitated as calcium salts, and the rapidly washed precipitate mixed with solution of potash, diluted, and filtered; the filtrate contains the tartrate, which is shown to be present by reprecipitation on boiling. The precipitate still on the filter is washed, dissolved in solution of chloride of ammonium, and the solution boiled; the citrate of calcium is reprecipitated. The presence of much sugar interferes with this reaction. A dilute solution of a citrate is not precipitated by chloride of calcium until the liquid is heated: precipitation from a strong solution, also, is not thoroughly complete without ebullition of the mixture. This reaction is not thoroughly satisfactory, citrate of calcium being *slightly* soluble in alkalies, in the solutions of salts produced in the reaction, and, to a very slight extent, even in cold water.

*Second Analytical Reaction.*—To a neutral solution of a citrate add solution of nitrate of silver; a white precipitate of citrate of silver ($Ag_3\overline{Ci}$) falls. Boil the mixture; the precipitate does not turn black as a tartrate of silver does, or only after long boiling.

*Third Analytical Reaction.*—To a neutral or alkaline solution of a citrate add a few drops of solution of permanganate of potassium and slowly heat the test-tube; reduction to manganate only occurs, a green or reddish-green solution resulting. Tartrates reduce the permanganate entirely.

*Other Analytical Reactions.*—Citric acid forms no precipitate corresponding with the acid tartrate of potassium.——Lime-water, in excess, gives no precipitate with citric acid or citrates, unless the solution is boiled, citrate of calcium being slightly soluble in cold but not in hot water; it usually precipitates tartrates in the cold.——Citrates, when heated with strong sulphuric acid, do not char immediately.——Citric acid and citrates prevent the precipitation of oxide of iron by alkalies, soluble double compounds being formed. The *Ferri et Ammo-*

*nii Citras*, U. S. P., is a preparation of this kind.——Metallic citrates decompose when heated, carbonates being formed and carbon set free: the odor of the gaseous products is not so characteristic as that of tartrates.

## QUESTIONS AND EXERCISES.

555. What is the source of citric acid?

556. Describe the method by which citric acid is prepared, giving diagrams

557. Illustrate by formulæ the various classes of tartrates and citrates.

558. State the average proportion of citric acid in lemon-juice.

559. Work out the sums proving the correctness of some of the figures given on page 325 as showing the saturating-power of citric acid for various carbonates.

560. What are the tests for citrates?

561. How are the tartrates separated from citrates?

## PHOSPHORIC ACID AND OTHER PHOSPHATES.

Formula of Phosphoric Acid $H_3PO_4$. Molecular weight 98.

*Source.*—The source of the ordinary normal phosphates and of phosphorus itself (*Phosphorus*, B. P. and U. S. P.) is the normal phosphate of calcium ($Ca_32PO_4$). It is the chief constituent of the bones of animals, being derived from the plants on which they feed, plants again obtaining it from the soil. Compounds of phosphorus are also met with in the brain, nerves, muscles, blood, saliva, and, according to Kirkes, even in tissues so simple that one must assume that the compounds are necessary constituents of the substance of the primary cell. They escape from the system both in the urine and fæces.

*Process.*—Phosphorus (P = 31) is obtained from bones by the following processes: The bones are burnt to remove all traces of animal matter. The resulting *bone-earth* is treated with sulphuric acid and water, by which an acid phosphate of calcium ($CaH_42PO_4$), often called *superphosphate of lime*, is produced:—

$$Ca_32PO_4 + 2H_2SO_4 = CaH_42PO_4 + 2CaSO_4.$$

The acid phosphate (strained from the sulphate and evaporated to dryness) is mixed with charcoal and sand, and heated to dull redness in an iron pot. At this stage water escapes and metaphosphate of calcium ($Ca2PO_3$, see Index) remains:—

$$CaH_42PO_4 = Ca2PO_3 + 2H_2O.$$

The mixture is then transferred to a retort and distilled at a strong red heat; silicate of calcium ($CaSiO_3$) is formed and remains in the

retort, phosphorus vapor is evolved and condensed under water, and carbonic oxide gas escapes:—

$$2(Ca2PO_3) + 2SiO_2 + C_{10} = 2CaSiO_3 + 10CO + P_4.$$

It is purified by melting under water containing sulphuric acid and red chromate of potassium.

*Properties.*—Phosphorus is a "semitransparent, colorless, wax-like solid (in sticks or cakes), which emits white vapors when exposed to the air. Specific gravity 1.77. It is soft and flexible at common temperatures, melts at 110° F., ignites in the air at a temperature a little above its melting-point, burning with a luminous flame and producing dense white fumes. It is very poisonous. Insoluble in water, but soluble in ether and in boiling oil of turpentine," also in bisulphide of carbon. It is soluble in oil which has been previously heated for a short time to about 300° F. to expel moisture; 12 grains in 4 fluidounces of dried almond oil constituting Phosphorated Oil, *Oleum Phosphoratum*, B. P. A mixture, or rather a solution, of phosphorus in balsam of tolu and yellow wax, forms the official Phosphorus Pill (*Pilula Phosphori*, B. P.).

*Granulated* or pulverulent phosphorus is obtained by placing a portion under equal parts of spirit and water in a bottle, standing the bottle in warm water till the phosphorus melts, then inserting the stopper (glass, not cork) and shaking the whole till cold.

*Red or Amorphous Phosphorus.*—Ordinary phosphorus kept at a temperature of about 450° Fahr., in an atmosphere from which air is excluded, becomes red, opaque, insoluble in liquids in which ordinary phosphorus is soluble, oxidizes extremely slowly, and only ignites when heated to near 500° Fahr. It is used in the manufacture of several varieties of lucifer-matches, not emitting the poisonous jaw-destroying fumes given by ordinary phosphorus.

*Quantivalence.*—The atom of phosphorus is quinquivalent, as seen in the pentachloride ($PCl_5$) and oxychloride ($PCl_3O$); but it often exhibits trivalent activity, as seen in the trichloride ($PCl_3$) and trihydride ($PH_3$).

*Molecular Weight.*—Phosphorus is an exception to the rule that the atomic weights (in grains, grammes, etc.) of elements occupy similar volumes of vapor at similar temperatures, the equivalent weight of phosphorus (31) only giving half such a volume. Hence while the molecular weights, that is, double the atomic weights, of oxygen ($O_2 = 32$), hydrogen ($H_2 = 2$), nitrogen ($N_2 = 28$), etc., give a similar bulk of vapor at any given temperature; the double atomic weight of phosphorus ($P_2 = 62$) only gives half this bulk; that is, four times the atomic weight of phosphorus must be taken to obtain the whole bulk. It would appear therefore that the molecule of phosphorus contains four atoms ($P_4 = 124$). As with sulphur, however, phosphorus in the state ordinarily known to us may be abnormal, and a variety yet be found in which the molecular weight is double the atomic weight.

## Phosphoric Acid.

The chief use of phosphorus in pharmacy is in the formation of Diluted Phosphoric Acid. Phosphorus is boiled with nitric acid and water until dissolved. The solution, evaporated to a low bulk to remove nitrous compounds, and rediluted so as to contain nearly 14 (13.8) per cent. of acid ($H_3PO_4$), equivalent to 10 per cent. of phosphoric anhydride ($P_2O_5$), constitutes the *Acidum Phosphoricum Dilutum*, B. P. and U. S. P., a colorless sour liquid of specific gravity 1.08 (1.056 U. S. P.). If the necessary appliances are at hand, four or five ounces of this acid may be prepared by the official process as follows: A quarter of a pint is made by boiling together, in a retort attached to a Leibig's condenser, 103 grains of phosphorus, 1½ fluidounces of the official nitric acid, and 2 ounces of water. When about 1 oz. of water has distilled over, it should be returned to the retort, and the operation repeated until the phosphorus has disappeared.

| $3P_4$ | + | $20HNO_3$ | + | $8H_2O$ | = | $12H_3PO_4$ | + | $20NO$ |
|---|---|---|---|---|---|---|---|---|
| Phosphorus. | | Nitric acid. | | Water. | | Phosphoric acid. | | Nitric oxide. |

The liquid remaining in the retort is then transferred to a dish (preferably of platinum), evaporated down to about half an ounce, and, lastly, diluted with distilled water to 5 fluidounces.

The use of the water in this process is to moderate the reaction. Strong hot nitric acid oxidizes phosphorus with almost explosive rapidity, hence the acid must be diluted in the first instance, and be rediluted, from time to time, to prevent its becoming too strong by loss of water. Time is saved by using a strong acid, but in that case constant supervision is necessary in order that water may be added, or the temperature otherwise reduced, when the action becomes too violent. Deficiency of nitric acid must also be avoided, or some phosphorous acid ($H_3PHO_3$) will be formed.

Markoe, also, to economize time, modifies the process by adding, for every ounce of phosphorus, four or five grains of iodine, and, drop by drop, twenty-five or thirty drops of bromine. The iodine and bromine unite with the phosphorus with a readiness or even violence that would be explosive if not controlled by the presence of the cold fluid—further cooled, if necessary, by immersing the vessel in cold water. Iodide of phosphorus ($PI_3$) and bromide of phosphorus ($PBr_3$) are at once formed. These, in the presence of water, immediately yield hydriodic and hydrobromic acids (HI, HBr) and phosphoric acid. The nitric acid attacks the hydriodic and hydrobromic acids, forming the lower oxides of nitrogen, which escape as gas, water, and free iodine and bromine. The latter unite with more phosphorus, and the reactions are repeated. This carrying power of a little iodine or bromine, or both, would perhaps be indefinitely pro-

longed if no vapor of these elements or their acids escaped with the gases. The phosphorus having disappeared, excess of nitric acid is got rid of roughly by dropping in clean rags or paper (nitric oxide, carbonic acid gas, and water being formed), and the last portions, by adding oxalic acid) which even still more readily yields similar products). Evaporation to a syrupy consistence finally removes all traces of iodine, bromine, oxalic acid, and moisture. The product is then diluted to any required extent.

Fig. 40.

*Experimental process.*—A flask, in the neck of which a funnel is inserted, and a second funnel inverted, so that its mouth rests within the mouth of the first, is an efficient and convenient arrangement of apparatus for this process, especially if the operation be conducted slowly.

Solution of phosphoric acid evaporated leaves a residue which melts at a low red heat, yielding *pyrophosphoric acid*, and, finally, *metaphosphoric acid* (*Glacial Phosphoric Acid*).

Phosphoric acid is also easily made from amorphous phosphorus (Mattison).

*Quantivalence.*—The elements represented by the formula $PO_4$ are those characteristic of phosphates. The grouping is trivalent; hence there may exist trimetallic or normal phosphates ($M'_3PO_4$), dimetallic acid phosphates ($M'_2HPO_4$), monometallic acid phosphates ($M'H_2PO_4$), and, lastly, trihydric phosphate ($H_3PO_4$) or common phosphoric acid. These are the ordinary phosphates met with in nature or used in pharmacy; the rarer pyrophosphates and metaphosphates, as well as the phosphites and hypophosphites, will be mentioned subsequently.

### *Analytical Reactions (Tests).*

***First Analytical Reaction.***—To an aqueous solution of a phosphate (*e. g.* $Na_2HPO_4$) add solution of sulphate of magnesium with which chloride of ammonium and ammonia have been mixed; a white crystalline precipitate of ammonio-magnesium phosphate falls ($MgAmPO_4$).

Chloride of ammonium is added to prevent the precipitation of hydrate of magnesium. Arseniates, which have close analogy to phosphites, give a similar precipitate with the magnesium reagent.

***Second Analytical Reaction.***—To a neutral aqueous solution of a phosphate add solution of nitrate of silver; light-yellow phosphate of silver ($Ag_3PO_4$) is precipitated. To a portion of the precipitate add ammonia; it dissolves. To another portion add nitric acid; it dissolves. By this reaction phosphates may

be distinguished from their close allies the arseniates, arseniate of silver being of a chocolate color.

*Third Analytical Reaction.*—To a solution (in a few drops of acid) of a phosphate insoluble in water (*e. g.* $Ca_2PO_4$) add the acetate of an alkali-metal (easily made by adding to soda or ammonia in a test-tube excess of acetic acid), and then a drop or two of solution of perchloride of iron; yellowish-white ferric phosphate ($Fe_2PO_4$) is precipitated. Too much of the ferric chloride must not be added, or ferric acetate will be produced, in which ferric phosphate is to some extent soluble.

*To remove the whole of the phosphoric radical* from the solution, add ferric chloride so long as a precipitate is produced, and boil; ferric phosphate and oxyacetate are precipitated.

*To obtain confirmatory evidence* of the presence of phosphate in this precipitate, and to separate the phosphoric radical as a pure unmixed phosphate, collect the precipitate on a filter, wash, drop some solution of ammonia on it, then sulphydrate of ammonium, and finally wash with water; black ferrous sulphide remains on the filter, while phosphate of ammonium occurs in the filtrate. To the filtrate add a mixture of solutions of sulphate of magnesium and chloride of ammonium, and well stir; ammonio-magnesian phosphate is precipitated.

*Fourth Analytical Reaction.*—To diluted nitric acid add a little phosphate of calcium (or any other phosphate) and then solution of molybdate of ammonium, and gently heat; a yellow precipitate falls.

This precipitate contains what is termed phospho-molybdic acid, but is a compound of molybdic acid with phosphoric acid (about 4 per cent. of $H_3PO_4$) and ammonia (nearly 7 per cent.).

Molybdate of ammonium is obtained by roasting the native sulphide of molybdenum ($MoS_2$) to molybdic oxide or anhydride ($MoO_3$), dissolving the latter in water, adding ammonia, evaporating, and crystallizing.

Molybdates having the following formulæ (M = 1 univalent atom of any metal) have been obtained: $M_2MoO_4$; $MHMoO_4$; $MHMoO_4$, $H_2MoO_4$. According to Carrington, commercial molybdate of ammonium is commonly the intermediate of these three salts.

*Note.*—The foregoing two reactions are useful in the analysis of bone-earth, other earthy phosphates, phosphate of iron, and all phosphates insoluble in water. Only arseniates give similar appearances; but the acid solution of these may be decomposed by agitation with sulphurous acid and subsequent treatment with sulphuretted hydrogen, arsenious sulphide, $As_2S_3$, being then precipitated.

*Other Analytical Reactions.*—Solutions of barium and calcium salts give, with aqueous solutions of phosphates, white

precipitates of the respective phosphates $BaHPO_4$, or $Ba_2PO_4$, and $CaHPO_4$, or $Ca_2PO_4$, all of which are soluble in acetic and the stronger acids.

## QUESTIONS AND EXERCISES.

562. State the source of phosphorus.
563. Give equations or diagrams explanatory of the isolation of phosphorus from its natural compounds.
564. What is the composition of farmers' "superphosphate," and how prepared?
565. Enumerate the properties of phosphorus.
566. Mention some solvents of phosphorus.
567. How is the official Diluted Phosphoric Acid made?
568. Describe the precautions necessary to be observed in making this acid.
569. What is the strength of the official acid?
570. Write formulæ illustrative of all classes of orthophosphates.
571. Mention the chief tests for soluble and insoluble phosphates.
572. By what reactions may phosphates be distinguished from arseniates?

*Vanadium*, V. 51.3, is a very rare element, and is here mentioned only because of its exceedingly interesting relationship to nitrogen, phosphorus, and arsenicum. Discovered but not isolated by Sefström, and its compounds investigated by Berzelius, it has only recently been obtained in the free state and fully studied by Roscoe.

| Oxides of Nitrogen. | | Oxides of Vanadium. | |
|---|---|---|---|
| $N_2O_5$, $N_2O_4$, $N_2O_3$, $N_2O_2$, $N_2O$. | | $V_2O_5$, $V_2O_4$, $V_2O_3$, $V_2O_2$, $V_2O$. | |
| Orthophosphates | $R_3'PO_4$ | Orthovanadates | $R_3'VO_4$ |
| Pyrophosphates | $R_4'P_2O_7$ | Pyrovanadates | $R_4'V_2O_7$ |
| Metaphosphates | $R'PO_3$ | Metavanadates | $R'VO_3$ |

Isomorphous Minerals.

| | |
|---|---|
| Apatite | $3(Ca_32PO_4),CaFl_2$ |
| Pyromorphite | $3(Pb_32PO_4),PbCl_2$ |
| Mimetesite | $3(Pb_32AsO_4),PbCl_2$ |
| Vanadinite | $3(Pb_32VO_4),PbCl_2$ |

## BORACIC ACID AND OTHER BORATES.

Formula of Boracic Acid $H_3BO_3$. Molecular weight 62.

The composition of artificial boracic acid is expressed by the formula $H_3BO_3$; but at a temperature of 212° F. this body loses the elements of water and yields metaboracic acid, $HBO_2$. The latter acid exists in the jets of steam (*fumerolles* or *suffioni*) that issue from the earth in some districts of Tuscany, and collects in the water of the *lagoni* (lagoons or little lakes) formed at the orifice of the steam-channel. This acid liquid, evaporated by aid of the waste natural steam and neutralized by carbonate of sodium, gives common borax ($2NaBO_2, 2HBO_2, 9H_2O$), possibly an acid metaborate of sodium with water of crystallization. It occurs "in transparent colorless crystals, sometimes slightly effloresced, with a weak alkaline reaction; insoluble in rectified spirit, soluble in water." Native borax, or *tincal*, and other borates are also found in Thibet, in Nevada, Peru, Chili, and recently in California, in the Clear Lake district. The introduction of the natural borax from California has reduced the price to about one-half its former amount. This borax is represented as forming large portions of the crystalline bed of a dried-up lake. Fused borax readily dissolves metallic oxides, as will have been already noticed in testing for cobalt and manganese. Hence, besides its use in medicine (*Sodii Boras*, U. S. P., *Borax*, *Mel Boracis*, B. P., or *Mel Sodii Boracis*, U. S. P., and *Glycerinum Boracis*, B. P.), it is employed as a flux in refining and other metallurgic and ceramic operations.

*Quantivalence.*—The boracic radical is trivalent ($BO_3'''$), the metaboracic, univalent ($BO_2'$); they have not been isolated. *The element boron*, like carbon, occurs in the amorphous, graphitoidal, and crystalline conditions. It is a trivalent element ($B'''$), yielding definite salts, such as the chloride ($BCl_3$) and fluoride ($BF_3$). Its atomic weight is 11.

### Reactions.

*First Synthetical Reaction.*—To a hot solution of a crystal of borax add a few drops of sulphuric acid and set aside; on cooling, crystalline scales of boracic acid ($H_3BO_3$) are deposited. They may be purified by collecting on a filter, slightly washing, drying, digesting in hot alcohol, filtering, and setting aside; pure boracic acid (B. P.) is deposited. The acid may also be recrystallized from water. Fifty grains dissolved in one ounce of rectified spirit constitute "Solution of Benzoic Acid," B. P.

When heated, these crystals lose water and yield, first, metaboracic acid ($HBO_2$), and, subsequently, when fused, boracic anhydride ($B_2O_3$).

Boracic acid is a very weak compound. It only slowly decom-

poses carbonates, and resembles alkaline substances in coloring turmeric brown. Indeed the alkalinity of borax is as great as if it contained no acidulous radical. Boracic acid is an antiseptic.

*Second Synthetical Reaction.*—Mix together 1 part of boracic acid, 4 of acid tartrate of potassium, and 10 or 20 of water; evaporate to a syrupy consistence, spread on plates, and set aside for dry scales to form. The resulting substance is, in water, far more readily soluble than either of its constituents, and is known as *boro-tartrate of potassium*, or *soluble cream of tartar*. The Prussian *tartarus boraxatus* differs from the foregoing French variety in containing 1 part of *borax* to 3 of acid tartrate of potassium.

### *Analytical Reactions (Tests).*

*First Analytical Reaction.*—Dip a piece of turmeric paper (paper soaked in tincture of turmeric tubers and dried) into a solution of boracic acid; it is colored brown-red, as by alkalies.

The usual way of applying this test is as follows: Add to a solution of any borate a few drops of hydrochloric acid, immerse half of a slip of turmeric paper in the liquid, then remove the hydrochloric acid by drying the paper over a flame. Concentrated hydrochloric acid and ferric chloride produce a somewhat similar effect.

*Second Analytical Reaction.*—To a fragment of a borate or metaborate (borax, for example) in a small dish or watch-glass add a drop of sulphuric acid and then a little alcohol, warm the mixture and set light to the spirit; the resulting flame will be tinged of a greenish color at its edges by the volatilized boracic acid or boracic anhydride.

The liquid should be well stirred while burning. Salts of copper and some metallic chlorides produce a somewhat similar color. The flame test may also be applied to a little of a mixture of the borax with strong sulphuric acid on a platinum wire. Glycerine may be used in place of sulphuric acid. (*Iles.*)

*Other Analytical Reactions.*—In solutions of borax barium salts give a white precipitate of barium metaborate ($Ba2BO_2$) soluble in acids and alkaline salts. Nitrate of silver gives metaborate of silver ($AgBO_2$) soluble in nitric acid and in ammonia. Chloride of calcium, if the solution is not too dilute, gives white borate of calcium.

## QUESTIONS AND EXERCISES.

573. Illustrate the relation of vanadium to nitrogen by formulæ of compounds of each element.

574. Describe the preparation of borax.

575. Give the formulæ of boracic acid, metaboracic acid, and borax.

576. Mention the tests for borates or metaborates.

---

*The foregoing acids and other salts contain the only acidulous radicals that are commonly met with in analysis or in ordinary medical or pharmaceutical operations. There are, however, many others which occasionally present themselves. The chief of these will now be shortly noticed; they are arranged in alphabetical order to facilitate reference.*

---

## SALTS OF RARER ACIDULOUS RADICALS.

BENZOIC ACID ($HC_7H_5O_2$) AND OTHER BENZOATES.—Slowly heat a fragment of benzoin (Gum benzoinum) (*Benzoinum*, B. P. and U. S. P.) in a test-tube; benzoic acid (*Acidum Benzoicum*, B. P. and U. S. P.) rises in vapor and condenses in small, white, feathery plates and needles, on the cool sides of the tube. If the benzoin is first mixed with twice its weight of sand or roughly powdered pumice-stone, and the heat very cautiously applied, the product will be less likely to be burnt, and a larger quantity be obtained. By repeated sublimation 10 to 15 per cent. may be obtained.

A more economical process is to boil the benzoin with one-fourth its weight of lime, filter, concentrate, decompose the solution of benzoate of calcium by hydrochloric acid, collect the precipitated benzoic acid, press between paper, dry and sublime in a tube or other vessel.

$$\underset{\text{Benzoic acid (impure).}}{2HC_7H_5O_2} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Benzoate of calcium.}}{Ca2C_7H_5O_2} + \underset{\text{Water.}}{2H_2O}$$

$$\underset{\text{Benzoate of calcium.}}{Ca2C_7H_5O_2} + \underset{\text{Hydrochloric acid.}}{2HCl} = \underset{\text{Chloride of calcium.}}{CaCl_2} + \underset{\text{Benzoic acid. (pure).}}{2HC_7H_5O_2}$$

There is always associated with the product a minute quantity of a volatile oil (styrol?) of agreeable odor, suggesting that of hay.

Benzoic acid is also prepared on a large scale artificially from naphthalene, one of the crystalline by-products in the distillation of

coal for gas. The naphthalene is oxidized by nitric acid to naphthalic or phthalic acid :—

$$\underset{\text{Naphthalene.}}{C_{10}H_8} + \underset{\text{Oxygen.}}{4O_2} = \underset{\text{Phthalic acid.}}{H_2C_8H_4O_4} + \underset{\text{Oxalic acid.}}{H_2C_2O_4}$$

The phthalic acid is neutralized by lime and the phthalate of calcium heated with hydrate of calcinm in a covered vessel at a temperature of about 640° F. for several hours. Benzoate and carbonate of calcium are formed, and, from the powder the benzoic acid is set free by action of hydrochloric acid.

$$\underset{\substack{\text{Phthalate of}\\\text{calcium.}}}{2CaC_8H_4O_4} + \underset{\substack{\text{Hydrate of}\\\text{calcium.}}}{Ca2HO} = \underset{\substack{\text{Benzoate of}\\\text{calcium}}}{Ca2C_7H_5O_2} + \underset{\substack{\text{Carbonate of}\\\text{calcium.}}}{2CaCO_3}$$

The crystalline deposit formed when essential oil of almonds (hydride of benzoyl or benzoic aldehyd) is exposed to the air is benzoic acid.

$$\underset{\substack{\text{Hydride of}\\\text{benzoyl.}}}{2C_7H_5OH} + \underset{\text{Oxygen.}}{O_2} = \underset{\substack{\text{Hydrate of}\\\text{benzoyl.}}}{2C_7H_5OHO} \text{ or } \underset{\substack{\text{Benzoic}\\\text{acid.}}}{2HC_7H_5O_2}$$

Benzoic acid is also obtained from hippuric acid (p. 338).

*Benzoate of Ammonium.*—To a little benzoic acid add a few drops of solution of ammonia; it readily dissolves, forming benzoate of ammonium (*Ammonii Benzoas*, U. S. P.) ($NH_4C_7H_5O_2$).

$$\underset{\text{Benzoic acid.}}{HC_7H_5O_2} + \underset{\text{Ammonia.}}{NH_4HO} = \underset{\substack{\text{Benzoate of}\\\text{ammonium.}}}{NH_4C_7H_5O_2} + \underset{\text{Water.}}{H_2O}$$

On evaporation, acid crystals or, ammonia being added, neutral crystals of benzoate of ammonium are deposited.

*Properties.*—Benzoic acid is also soluble in other alkaline liquids, forming benzoates. It is slightly soluble in cold water, more so in hot, and readily soluble in rectified spirits. It melts at 248° F., and boils at 462°, volatilizing with only a slight residue.

*Tests.*—The following are the *tests* for benzoic acid: To a portion of the above solution of benzoate of ammonium add a drop or two of sulphuric or hydrochloric acid; a white crystalline precipitate of benzoic acid separates. To another portion, carefully made neutral, add a drop or two of neutral solution of perchloride of iron; reddish ferric benzoate is precipitated.

*Cinnamic Acid* ($C_9H_8O_2$).—Benzoic acid is distinguished from an allied body, cinnamic acid (occurring in Balsams of Peru, Tolu, and Storax, and sometimes in Benzoin), by not yielding hydride of benzoyl ($C_7H_5OH$) (oil of bitter almonds) when distilled with chromic acid—that is, with a mixture of red chromate of potassium and sulphuric acid.

CARMINIC ACID ($C_{14}H_{14}O_8$).—This is the coloring principle of cochineal (*Coccus*, B. P. and U. S. P.). The *carmine* of trade, when unadulterated (*vide* Pharmaceutical Journal, 1859–60, p. 546), is carminic acid united with one or two per cent. of alumina, or, occasionally, of oxide of tin or albumen. It should be wholly soluble in solution of ammonia, giving an apparently clear rich purple fluid. Carmine, with French chalk, or starch, constitutes *face rouge* or *animal rouge*.

Merrick tests the relative value of several samples of cochineal or carmine by observing how much solution of permanganate of potassium is required to change the color of a decoction to faint pink.

CETRARIC ACID ($H_2C_{24}H_{30}O_{16}$) is the bitter principle of Iceland moss (*Cetraria*, B. P. and U. S. P.). In the lichen it is associated with much starch.

CHRYSOPHANIC ACID ($C_{14}H_8O_4$?).—This acid is the chief coloring-matter of various species of rhubarb root (*Rhei Radix*, B. P., *Rheum*, U. S. P.), and, under the name of *parietinic acid*, of various common yellow lichens. The author also finds it to form four-fifths of "Chrysarobine," a name given by Kemp to the pith, etc., of a leguminous tree. Chrysarobine is also known as Araroba Powder, Bahia Powder, Brazil Powder, Goa Powder, and Ringworm Powder. Chrysophanic acid may be obtained in crystals of a golden-yellow color, hence the name (from χρυσὸς, *chrusos*, gold, and φαίνω, *phainō*, I shine). Its synonyms are *Rhaponticin*, *Rheic acid*, *Rhein*, *Rheumin*, *Rheubarbaric acid*, *Rheubarbarin*, *Rumicin*. Chrysophanic acid, black, red-brown, and red resins (*Aporetine*, *Phœoretine*, and *Erythroretine*), a bitter principle, and tannic acid are considered to be the conjoint source of the therapeutic properties of rhubarb. Chrysophanic acid is also contained in several species of *Rumex* or *Dock*. "Rumicin" is a preparation of Yellow Dock.

CYANIC ACID (HCyO) AND OTHER CYANATES.—The valuable reducing power of cyanide of potassium (KCy) (or ferrocyanide, $K_4Fcy$) on metallic compounds is due to the avidity with which cyanate (KCyO) is formed.

*Process.*—Fuse a few grains of cyanide of potassium in a small porcelain crucible, and add powdered oxide of lead; a globule of metallic lead is at once set free, excess of the oxide converting the whole of the cyanide of potassium into cyanate of potassium.

*Urea.*—Cyanate of potassium (KCNO), or, better, cyanate of lead (Pb2CNO), treated with sulphate of ammonium, yields cyanate of ammonium ($NH_4CNO$); and solution of cyanate of ammonium, when simply heated, changes to artificial *urea* ($CH_4N_2O$), the most important constituent of urine, and the chief form in which the nitrogen of food is eliminated from the animal system. The process will be more fully described subsequently in connection with urea.

FORMIC ACID ($HCHO_2$).—The red ant (Formica rufa) and several other insects, when irritated, eject a strongly acid, acrid liquid,

having a composition expressed by the above formula, and which has appropriately received the name of formic acid; it is also contained in the leaves of the stinging-nettle.

*Process.*—It may be artificially prepared by heating equal weights of oxalic acid and glycerine to a temperature of from 212° F. to 220° for fifteen hours. The glycerine has, apparently, no chemical action, but, for some unknown reason, induces decomposition of the oxalic acid at a lower temperature than would otherwise be necessary; at a higher temperature the formic acid itself is decomposed. On distilling the mixture with water, the formic acid slowly passes over. The dilute acid may be concentrated by neutralizing with carbonate of lead, filtering, evaporating to a small bulk, collecting the deposited crystalline formate of lead, drying, decomposing in a current of sulphuretted hydrogen, and separating the resulting syrupy acid, or distilling the formate of lead with strong sulphuric acid.

$$\underset{\text{Oxalic acid.}}{H_2C_2O_4} = \underset{\text{Formic acid.}}{HCHO_2} + \underset{\text{Carbonic acid gas.}}{CO_2}$$

*Formic acid* may be instructively though not economically prepared by the oxidation of methylic alcohol (wood-spirit), just as acetic acid and valerianic acid are obtained from ethylic alcohol and amylic alcohol respectively.

$$\underset{\text{Wood spirit.}}{CH_3HO} + \underset{\text{Oxygen.}}{O_2} = \underset{\text{Formic acid.}}{HCHO_2} + \underset{\text{Water.}}{H_2O}$$

*Tests.*—Formic acid does not char when heated alone or with sulphuric acid, but splits up into carbonic oxide gas and water. It is recognized by this property and by its reducing action on salts of gold, platinum, mercury, and silver. It is solid below 32° F.

GALLIC ACID.—See *Tannic Acid.*

HEMIDESMIC ACID.—The supposed active principle of hemidesmus root (*Hemidesmi Radix*, B. P.).

HIPPURIC ACID ($HC_9H_8NO_3$) is a constituent of human urine (much increased on taking benzoic acid), but is best prepared from the urine of the horse (hence the name, from ἵππος, *hippos*, a horse), or, better, from that of the cow. To such urine add a little milk of lime, boil for a few minutes, remove precipitated phosphates by filtration, drop in hydrochloric acid until the liquid, after well stirring, is exactly neutral to test-paper, concentrate to about one-eighth the original bulk, and add excess of strong hydrochloric acid; impure hippuric acid is deposited. From a solution of the impure acid in hot water chlorine gas removes the color, and the liquid deposits crystals of pure hippuric acid on cooling. Its constitution is that of *benzoic glycocine*, $C_2H_2H_2(C_7H_5O)NO_2$.

*Tests.*—To a solution of hippurate add neutral solution of ferric chloride; a brown precipitate (ferric hippurate) results. Salts of silver and mercury give white precipitates. Heat hippuric acid in a test-tube; it chars, benzoic acid sublimes, and vapors of characteristic odor are evolved; they contain, amongst other bodies, hydrocyanic acid, and a substance smelling somewhat like Tonka bean.——The crystalline form of hippuric acid is characteristic; it will be described in connection with the subject of urine.

## QUESTIONS AND EXERCISES.

577. Give the preparation, composition, properties, and tests of benzoic acid, employing equations or diagrams.
578. What is the nature of carmine?
579. Name the bitter principle of Iceland moss.
580. Mention the coloring principle of rhubarb.
581. To what is rhubarb considered to owe its medicinal activity?
582. How is cyanate of potassium prepared, how converted into an ammonium salt, and what are the relations of the latter to urea?
583. Give the formulæ of cyanic acid, cyanate of ammonium, and urea.
584. What is the chemical formula of formic acid?
585. Describe the artificial production of formic acid.
586. Describe the relation of formic acid to wood-spirit.
587. State the sources, characters, and tests of hippuric acid.

HYDROFERROCYANIC ACID ($H_4Fe''Cy_6$, or $H_4Fcy''''$) AND OTHER FERROCYANIDES.—The ferrocyanide of most interest is that of potassium (*Potassii Ferrocyanidum*, U. S. P.), the yellow prussiate of potash (*Potassæ Prussias Flava*, B. P.) ($K_4FeC_6N_6,3H_2O$), the formation of which was alluded to in connection with hydrocyanic acid (see page 279). It cannot be regarded as simply a double salt of cyanide of potassium with cyanide of iron ($FeCy_2,4KCy$), its chemical properties being entirely different from either of those substances; moreover, unlike cyanide of potassium, it is not poisonous. Most of its reactions point to the conclusion that its iron and cyanogen are intimately united to form a definite quadrivalent radical appropriately termed *ferrocyanogen* ($FeCy_6$, or Fcy). One part of ferrocyanide of potassium in 20 of water forms the official "Solution of Yellow Prussiate of Potash," B. P.

*Tests.*—Many of the ferrocyanides are insoluble, and are therefore precipitated when solution of ferrocyanide of potassium is added to the various salts. Those of iron and copper, being of characteristic color, are adopted as tests of the presence of the metals or of the ferrocyanogen, as the case may be.

To solution of ferrocyanide of potassium add a ferric salt;

ferrocyanide of iron ($Fe_4Fcy_3$) (Prussian blue) (*Ferri ferrocyanidum*, U. S. P.) is precipitated.

$$3K_4Fcy + 2(Fe_2 3SO_4) = Fe_4Fcy_3 + 6K_2SO_4.$$

To another portion add solution of a copper salt; reddish-brown ferrocyanide of copper ($Cu_2Fcy$) is precipitated.

*Note.*—The ferrocyanogen in ferrocyanide of potassium is broken up when the salt is heated with sulphuric acid, *carbonic oxide* being evolved if the acid is strong (that is, ordinary oil of vitriol—$H_2SO_4$ with 3 to 4 per cent. of water), and *hydrocyanic acid* if weak:—

$$K_4FeC_6N_6, 3H_2O + 3H_2O + 6H_2SO_4 = 2K_2SO_4 + FeSO_4 + 3(NH_4)_2SO_4 + 6CO.$$

$$2K_4FeCy_6 + 6H_2SO_4 + xH_2O = FeK_2FeCy_6 + 6KHSO_4 + 6HCy + xH_2O.$$

*Hydrocyanic Acid* has already been described. (*Vide* p. 280.)

*Carbonic Oxide* (CO).—Heat two or three fragments of ferrocyanide of potassium with eight or ten times their weight of sulphuric acid, and as soon as the gas begins to be evolved, remove the test-tube from the flame; for the action, when once set up, proceeds somewhat tumultuously. Ignite the carbonic oxide at the mouth of the tube; it burns with a pale blue flame, the product of combustion being carbonic acid gas ($CO_2$).

Carbonic oxide is a direct poison. It is generated whenever coke, charcoal, or coal burns with an insufficient supply of air. Hence the danger of open fires in the more or less closed apartments of ordinary dwellings.

Carbonic oxide may also be obtained from oxalic acid. (*Vide* p. 317.)

Hydroferridcyanic Acid ($H_6Fe'''_2Cy_{12}$, or $H^I_6Fdcy^{VI}$) and other Ferridcyanides.—Pass chlorine gas slowly through solution of ferrocyanide of potassium until the liquid, after frequent shaking, ceases to give a blue precipitate, when a minute portion is taken out on the end of a glass rod and brought into contact with a drop of a dilute solution of a ferric salt; it now contains ferridcyanide of potassium ($K_6Fe'''_2Cy_{12}$, or $K^I_6Fdcy^{VI}$), *red prussiate of potash* (B. P.), as it is termed from the color of its crystals. Excess of chlorine must be carefully avoided, as chloride of cyanogen and other compounds are then formed.

$$2K'_4Fe''Cy'_6 + Cl'_2 = 2K'Cl' + K'_6Fe'''_2Cy'_{12}.$$

*Another process.*—To a cold solution of yellow prussiate of potash so much hydrochloric acid is added as will take two atoms of potassium from two molecules of the salt, and then a cold clear solution of bleaching-powder till ferric chloride gives

no reaction. Any excess of acid is then neutralized with chalk and the solution evaporated to crystallization (Rhien).

*Note.*—The removal of two atoms of potassium from the ferrocyanide is the only change of composition that occurs; but the ferrocyanogen is altered in quality, its iron passing from the ferrous to the ferric condition, from bivalent to trivalent activity, altered to a condition in which it no longer precipitates ferric salts, but, on the other hand, gives a dark-blue precipitate with ferrous salts. The radical is distinguished as ferridcyanogen.

Ferridcyanide of potassium may also be prepared by a modification of the foregoing method in which nascent instead of free chlorine is employed (Wenzell). Take of Bichromate of Potassium 1 part, Ferrocyanide of Potassium Cryst. 5.72 parts, Hydrochloric Acid, of spec. grav. 1.16, 3 parts, by weight, Water 60 parts. Dissolve the two salts in hot water, add the acid, heat to boiling, continuing the ebullition, replacing the water evaporated during the process until a portion of the filtered liquid is not precipitated on the addition of sesquichloride of iron. When reaction is completed, filter the liquid and wash the hydrate of chromium, unite the liquids, and concentrate to crystallization. If the evaporated liquid possess an acid reaction, the addition of caustic potash, in sufficient quantity to cause a weak alkaline reaction, will greatly facilitate the subsequent crystallization.

$$6(K_4FeCy_6) + K_2Cr_2O_7 + 8HCl = 3(K_6Fe_2Cy_{12}) + 8KCl + H_2O + Cr_26HO.$$

*Test.*—To a portion of the solution add solution of ferrous sulphate; a precipitate falls. This precipitate is ferridcyanide of iron (Turnbull's blue), $Fe''_3Fe'''_2Cy'_{12}$, or $Fe^{II}_3Fdcy^{VI}$.

$$K_6Fdcy + 3FeSO_4 = Fe_3Fdcy + 3K_2SO_4.$$

It will be noticed that the change in the condition of the iron keeps up the balance of the atomic values of the various parts of the radicals or of the salts; the quantivalential equilibrium is maintained

A solution of 1 of ferridcyanide of potassium in 20 of water constitutes the "Solution of Red Prussiate of Potash," B. P.

HYDROFLUORIC ACID (HF) AND OTHER FLUORIDES.—Molecular weight of HF, 20. The chief use of hydrofluoric acid is in the etching on glass. The operation, performed on the small scale, also constitutes the best test for fluorine, the elementary radical of all fluorides.

*Process and Test.*—Warm any odd piece of window-glass, having an inch or two of surface, until a piece of beeswax rubbed on one side yields a thin oily film. When cool make a cross, letter, or other mark on the glass by pressing a pointed piece of wood, a penknife, or file, through the wax. Place a few grains of powdered fluor spar, the commonest natural fluor-

ide, in a porcelain crucible (or a lead cup), add a drop or two of sulphuric acid, cover the crucible with the prepared glass, waxed side downwards, and gently warm the bottom of the crucible in a fume-chamber or in the open air, in such a way as not to melt the wax. After a few minutes remove the glass, wash the waxed side by pouring water over it, scrape off most of the wax, then warm the glass, and wipe off the remainder; the marks made through the wax will be found to be permanently etched on the glass; the acid has eaten into or *etched* (from the German *ätzen*, to corrode) the glass.

In the above operation the fluoride of calcium and sulphuric acid yield hydrofluoric acid, thus:—

$$CaF_2 + H_2SO_4 = CaSO_4 + 2HF.$$

The hydrofluoric acid gas and the silica of the glass then yield gaseous fluoride of silicon ($SiF_4$), which escapes, and water, thus:—

$$4HF + SiO_2 = 2H_2O + SiF_4.$$

The silica, being removed from the glass, leaves furrows or etched portions.

*Note.*—In the experiment just described, the liberated hydrofluoric acid also attacks the siliceous glazing of the porcelain crucible; so that in important cases, where search is made for very small quantities of fluorine, vessels of platinum or lead must be employed.

*Uses.*—The aqueous solution of hydrofluoric acid used by etchers, and commonly termed simply hydrofluoric acid, or fluoric acid, is prepared in leaden stills and receivers, and kept in leaden or gutta-percha bottles. Except these materials, as well as platinum and fluor spar, hydrofluoric acid rapidly attacks any substance of which bottles and basins are usually made. It quickly cauterizes the skin, producing a painful slow-healing sore.

*Quantivalence.*—The atom of fluorine, like that of chlorine, bromine, or iodine, is univalent (F′). The great analogy existing between these radicals extends to their compounds.

Fluorine is said to be a colorless gas; but, from the avidity with which it combines with all elements (except oxygen), it is so difficult of isolation as hitherto to preclude satisfactory study of its physical properties.

HYPOPHOSPHOROUS ACID ($H_3PO_2$, or $HPH_2O_2$) AND OTHER HYPOPHOSPHITES.—Boil together, in a fume-chamber, a grain or two of phosphorus, a few grains of slaked lime, and about a quarter of an ounce of water until phosphoretted hydrogen, a spontaneously inflammable, badly smelling gas, ceases to be evolved. The mixture, filtered, and excess of lime removed by carbonic acid gas, yields solution of hypophosphite of calcium ($Ca2PH_2O_2$)—(*Calcis Hypophosphis*, B. P.).

$$P_8 + 6H_2O + 3CaH_2O_2 = 3(Ca2PH_2O_2) + 2PH_3.$$

The solution, when concentrated by evaporation, has been known to explode, probably from formation of phosphoretted hydrogen. This may be prevented, it is said, by evaporating at a low temperature, especially towards the close of the operation; or by adding alcohol, which decomposes any traces of liquid phosphoretted hydrogen ($PH_2$) or solid phosphoretted hydrogen ($P_2H$) which possibly may be present, and to which it is conceivable explosion may be due.

*Phosphoretted hydrogen* ($PH_3$).—The above reaction is also that by which phosphoretted hydrogen, the third hydride of phosphorus, may be prepared. If the gas is to be collected, the phosphorus and water may first be boiled in a flask until a jet of spontaneously inflammable phosphorus vapor escapes, with steam, from the end of the attached delivery-tube. Strong solution of caustic potash or soda is next very gradually poured into the flask through a funnel tube previously fitted into the cork, the liquid being kept boiling. Phosphoretted hydrogen is then evolved, and, if the delivery-tube dip under water, may be collected, or allowed to slowly pass up through the water bubble by bubble so as to form the peculiar rings of smoke (phosphoric anhydride) characteristic of the experiment.

*Hypophosphite of calcium* (*Calcii Hypophosphis*, U. S. P.) may be obtained in crystals by evaporating and slowly cooling; but the solution is usually at once evaporated to dryness, a white pulverulent salt being obtained. The hypophosphites of Potassium and Sodium (*Potassii Hypophosphis*, U. S. P., and *Sodii Hypophosphis*, U. S. P.) may be obtained in the same way from their hydrates, and many other hypophosphites similarly from other hydrates, or by double decomposition of the calcium salt and carbonates.

*Hypophosphite of sodium* ($NaPH_2O_2, H_2O$) (*Sodæ Hypophosphis*, B. P.) is made by decomposing solution of hypophosphite of calcium by carbonate of sodium, filtering, and evaporating to dryness. It is a white, granular, deliquescent substance.

$$Ca2PH_2O_2 + Na_2CO_3 = 2NaPH_2O_2 + CaCO_3.$$

When heated, the water is first evolved, then hydrogen and spontaneously inflammable phosphoretted hydrogen, and a mixture of pyrophosphate and metaphosphate of sodium remains (Rammelsberg).

$$5NaPH_2O_2 = Na_4P_2O_7 + NaPO_3 + 2PH_3 + 2H_2.$$

*Hypophosphorous acid*, the hydrogen hypophosphite, may be prepared by decomposing the barium salt with sulphuric acid or the calcium salt by oxalic acid; hypophosphite of quinine by dissolving the alkaloid in hypophosphorous acid, or by decomposing sulphate of quinine by hypophosphite of barium. Hypophosphite of Iron (*Ferri Hypophosphis*, U. S. P.) may be obtained by dissolving ferric hydrate in cold aqueous hypophosphorous acid and evaporating the solution.

The hypophosphites are often used in medicine in the form of syrups. The term hypophosphite is in allusion to the smaller amount (*ὑπὸ*, *hupo*, under or deficiency) of oxygen in these com-

pounds ($R'_3PO_2$) than in the phosphites ($R_2PO_3$), a class of salts having again less oxygen in their molecules than exists in those of the phosphates ($R_2PO_4$). The prefix *hypo* has similar significance in such words as hyposulphite and hypochlorite.

*Tests.*—To a portion of the above solution of hypophosphite of calcium add solution of chloride of barium, chloride of calcium, or acetate of lead; in neither case is a precipitate obtained, whereas soluble phosphates and phosphites yield white precipitates of phosphate or phosphite of barium, calcium, or lead. To other portions add solutions of nitrate of silver and mercuric chloride; the respective metals are precipitated as by phosphites. To another small portion add zinc and dilute sulphuric acid; hydrogen and phosphoretted hydrogen are evolved as from phosphites. To another portion add sufficient oxalic acid to remove the calcium; filter; to the solution of hypophosphorous acid add solution of sulphate of copper, and slowly warm the mixture; solid brown cuprous hydride ($Cu_2H_2$) is precipitated: increase the heat to the boiling point; hydrogen is evolved and metallic copper set free. Heat a small quantity of a solid hypophosphite on the end of a spatula in a flame; it splits up into pyrophosphate, a little metaphosphate, hydrogen, phosphoretted hydrogen, and water, burning with a phosphorescent light—the official hypophosphite of calcium yielding about 80 per cent. of residue.

$$7(Ca2PH_2O_2) = 3Ca_2P_2O_7 + Ca2PO_3 + 6PH_3 + H_2O + 4H_2.$$

To an acid solution of a hypophosphite add solution of permanganate of potassium. It is instantly decolorized in the cold. The same effect follows its addition to acid solution of a phosphite, but not to that of an ortho-, meta-, or pyrophosphate.

HYPOSULPHUROUS ACID ($H_2S_2O_2$) AND OTHER HYPOSULPHITES.—The only hyposulphite of much interest in pharmacy is the sodium salt (Hyposulphite of Soda, B. P., *Sodii Hyposulphis*, U. S. P.) ($Na_2S_2O_3,5H_2O$).

*Process.*—Heat together gently, or set aside in a warm place, a mixture of solution of sulphite of sodium ($Na_2SO_3$), and a little powdered sulphur; combination slowly takes place, and hyposulphite of sodium is formed. The solution, filtered from excess of sulphur, readily yields crystals. [The solution of sulphite of sodium may be made by saturating a solution of soda with sulphurous acid gas.]

*Use of hyposulphite of sodium in quantitative analysis.*—In the British Pharmacopœia hyposulphite of sodium is given

as a reagent for the quantitative estimation of free iodine in volumetric analysis. To a few drops of iodine-water add cold mucilage of starch; a deep-blue color (starch iodide) is produced. To the product add solution of hyposulphite of sodium until the blue color just disappears. This absorption of iodine is sufficiently definite and delicate to admit of application for quantitative purposes. It depends on the combination of the iodine with half of the sodium in two molecules of the hyposulphite, the hyposulphurous radicals of the two molecules apparently coalescing to form a new radical, the tetrathionic (from τέτρας, *tetras*, four, and θεῖον, *theion*, sulphur), tetrathionate of sodium ($Na_2S_4O_6$) and iodide of sodium being formed.

*Sulphur Oxyacids.*—It will be as well here to give the formulæ of other oxyacids of sulphur, forming with the four already mentioned a series that is as useful as the series of compounds of nitrogen and oxygen in illustrating the soundness of Dalton's atomic theory (p. 50).

| | |
|---|---|
| Hydrosulphurous Acid . . . . . | $H_2SO_2$ |
| Sulphurous Acid . . . . . | $H_2SO_3$ |
| Sulphuric Acid . . . . . | $H_2SO_4$ |
| Hyposulphurous Acid . . . . . | $H_2S_2O_2$ |
| Dithionic Acid . . . . . | $H_2S_2O_6$ |
| Trithionic Acid . . . . . | $H_2S_3O_6$ |
| Tetrathionic Acid . . . . . | $H_2S_4O_6$ |
| Pentathionic Acid . . . . . | $H_2S_5O_6$ |

*Use of "Hypo" in Photography.*—The sodium hyposulphite is largely used in photography to dissolve chloride, bromide, or iodide of silver off plates which have been exposed in the camera. Prepare a little chloride of silver by adding a chloride (chloride of sodium) to a few drops of solution of nitrate of silver. Collect the precipitated chloride on a filter, wash, and add a few drops of solution of hypophosphide of sodium; the silver salt is dissolved, solution of double hyposulphite of sodium and silver being formed. The solution of this double hyposulphite has a remarkably sweet taste, sweeter than syrup, if the solution is strong. The double hyposulphite of sodium and gold is employed for giving a pleasant tint to photographic prints.

*Test.*—To solution of a hyposulphite add a few drops of dilute sulphuric or other acid; hyposulphurous acid is set free, but at once begins to decompose into sulphurous acid, recognized by its odor, and free sulphur ($2H_2S_2O_2 = 2H_2SO_3 + S_2$). This reaction constitutes the best test for hyposulphites. Another good test of a soluble simple hyposulphite is its power of dissolving chloride of silver with production of a more or less sweet solution.

## QUESTIONS AND EXERCISES.

588. Give the formula of ferrocyanide of potassium.
589. What is the supposed constitution of ferrocyanide of potassium?
590. Enumerate the tests for ferrocyanogen.
591. What are the respective reactions of ferrocyanide of potassium with strong and weak sulphuric acid?
592. Mention and explain a common source of carbonic oxide in households. What is the product of its combustion?
593. Write equations or diagrams illustrative of the changes effected on ferrocyanide of potassium during its conversion into ferridcyanide.
594. By what reactions may the presence of a ferridcyanide in a solution be demonstrated?
595. State the difference between Prussian blue and Turnbull's blue.
596. Describe the source, mode of preparation, chief use of, and test for hydrofluoric acid.
597. Illustrate by a diagram the preparation and composition of hyposulphite of sodium.
598. Mention the uses and characteristic reactions of hyposulphite of sodium.
599. Give the names and formulæ of eight acids, each containing hydrogen, sulphur, and oxygen.

---

LACTIC ACID ($H_3C_3H_4O_3$) AND OTHER LACTATES.—Lactic acid occurs naturally in willow bark (Dott). When milk turns sour its sugar has become converted into an acid appropriately termed lactic (*lac*, *lactis*). Other saccharine and amylaceous substances also by fermentation yield lactic acid. Neither the hydrogen lactate (lactic acid) nor other lactates are much used in England, but the former is official in America (*Acidum Lacticum*, U. S. P.).

*Process.*—Lactate of calcium and lactic acid may be prepared ar follows: Mix together eight parts of sugar, one of common cheese, three of chalk, and fifty of water, and set aside in a warm place (about 80° F.) for two or three weeks; a mass of small crystals of lactate of calcium results. Remove these, recrystallize from hot water, decompose by sulphuric acid, avoiding excess, digest in alcohol, filter off the sulphate of calcium, evaporate the clear solution to a syrup; this residue is lactic acid; sp. gr. 1.212.

*Lactate of Iron* (*Ferri Lactas*, U. S. P.) is made by digesting iron filings in warm diluted lactic acid (1 acid to 16 water) till effervescence of hydrogen ceases, filtering and setting aside to cool and crystallize. The crystals are collected, washed with alcohol, and dried. This ferrous lactate occurs in greenish-

white crystalline crusts or grains, of a mild, sweetish, ferruginous taste, soluble in forty-eight parts of cold and twelve of boiling water, but insoluble in alcohol. Exposed to heat it froths up, gives out thick, white, acid fumes, and becomes black; sesquioxide of iron being left. If it be boiled for fifteen minutes with nitric acid of the specifie gravity 1.20, a white, granular deposit of mucic acid will occur on the cooling of the liquid.

*Tests.*—No single reaction of lactic acid is sufficiently distinctive to form a test. The crystalline form of the lactate of calcium, as seen by the microscope, is characteristic. The production of this salt, and the isolation of the syrupy acid itself, are the only means, short of quantitative analysis, on which reliance can be placed.

*A variety of lactic acid* has been obtained from the juice of fish; it is termed *sarcolactic* acid (from σάρξ, σαρκὸς *sarx, sarcos*, flesh).

MALIC ACID ($H_2C_4H_2O_5$) AND OTHER MALATES (from *malum*, an apple).—The juice of unripe apples, gooseberries, currants, rhubarb stalks, etc., contains malic acid and malate of potassium. When isolated it occurs in deliquescent prismatic crystals.

*Tests.*—Malate of calcium ($CaHC_4H_3O_5$) is soluble in water; hence the aqueous solution of malic acid or other malate is not precipitated by lime-water or chloride of calcium; but on adding spirit of wine a white precipitate falls, owing to the insolubility of the calcium malate in alcohol. Malates are precipitated by lead-salts; on warming the malate of lead with acetic acid it dissolves, separating out in acicular crystals on cooling. If the mixture be heated without acid, the malate of lead agglutinates and fuses.

Hot strong sulphuric acid chars malic acid far less readily than it does nearly all other organic acids.

*Asparagin* ($C_4H_8N_2O_3,H_2O$).—This proximate principle of plants occurs in many vegetable juices, and doubtless plays a very important part in their nutrition. It is deposited in crystals when the fresh juices of asparagus, marsh-mallow, etc., are rapidly evaporated. It is noticed here because malic acid is readily obtained from it by oxidation, nitrogen being eliminated, and because its exact natural position among chemical substances is not yet well made out. The atoms of its molecule are those of aspartate of ammonium ($NH_4C_4H_6NO_4$), into which it is converted when its solution is long boiled. Decomposed by aid of ferments, asparagin, absorbing hydrogen, yields succinate of ammonium $(NH_4)_2C_4H_4O_4$. Such reactions as these and the formation of lactic acid from sugars should suggest to the student possible modes in which chemical changes take place in the plant-department of the vast laboratory of nature.

MECONIC ACID ($H_3C_7HO_7$).—Opium contains meconic acid (from *μήκων, mekōn*, a poppy) partially combined with morphia. To concentrated infusion of opium nearly neutralized by ammonia, add solution of chloride of calcium; meconate of calcium is precipitated. Wash the precipitate, place it in a small quantity of hot water, and add a little hydrochloric acid; the clear liquid (filtered, if necessary) deposits scales of meconic acid on cooling.

*Tests.*—To solution of meconic acid or other meconate, or to infusion of opium, add a neutral solution of ferric chloride; a red solution of meconate of iron is produced. To a portion of the mixture add solution of corrosive sublimate; the color is not destroyed; to another portion add hydrochloric acid; the color is discharged. (These reagents act on sulphocyanate of iron, which is of similar tint, in exactly the opposite manner.)

The normal meconates of potassium, sodium, and ammonium are soluble in water, the acid meconates very slightly soluble, the meconates of barium, calcium, lead, copper, and silver insoluble in water but soluble in acetic acid.

METAPHOSPHORIC ACID ($HPO_3$) AND OTHER METAPHOSPHATES.—Prepare phosphoric anhydride ($P_2O_5$) by burning a small piece of phosphorus in a porcelain crucible placed on a plate and covered by an inverted test-glass tumbler, half-pint measure-glass, or some such vessel. After waiting a few minutes for the phosphoric anhydride to fall, pour a little water on the plate and filter the liquid; the product is solution of metaphosphoric acid (from *μετά, meta*, a preposition denoting change).

$$P_2O_5 + H_2O = 2HPO_3.$$

*Tests.*—To solution of metaphosphoric acid add ammonio-nitrate of silver, or to a neutral metaphosphate add solution of nitrate of silver; a white precipitate ($AgPO_3$) is obtained. This reaction sufficiently distinguishes metaphosphates from the ordinary phosphates or orthophosphates (from *ὀρθός, orthos*, straight), as the common phosphates may, for distinction, be termed (which give, it will be remembered, a *yellow* precipitate with nitrate of silver). Another variety of phosphates shortly to be considered, the pyrophosphates, also gives a white precipitate with nitrate of silver. To the solution of metaphosphoric acid obtained as above or by the action of acetic acid on a metaphosphate, add an aqueous solution of white of egg; coagulation of the albumen ensues. Neither orthophosphoric nor pyrophosphoric acid coagulates albumen. Boil the aqueous solution of metaphosphoric acid for some time; on testing the

solution the acid will be found to have been converted into orthophosphoric acid:—

$$HPO_3 + H_2O = H_3PO_4 \text{ (orthophosphoric acid).}$$

The ordinary medicinal phosphoric acid is made from phosphorus and nitric acid, the liquid being evaporated to a syrupy consistence to remove the last traces of nitric acid. It may contain pyrophosphoric and metaphosphoric acids, if the heat employed be high enough to remove the elements of water:—

$$2H_3PO_4 - H_2O = H_4P_2O_7 \text{ (pyrophosphoric acid).}$$
$$H_3PO_4 - H_2O = HPO_3 \text{ (metaphosphoric acid).}$$

On redilution the metaphosphoric acid only slowly reabsorbs water. If, therefore, on testing, metaphosphoric be found to be present, the solution should be boiled until conversion to orthophosphoric acid has occurred.

NITROUS ACID ($HNO_2$) AND OTHER NITRITES.—Strongly heat a fragment of nitrate of potassium or sodium on a piece of platinum foil; oxygen is evolved and nitrite of potassium remains.

*Test.*—Dissolve the residue in water, add a few drops of dilute sulphuric acid, then a little weak solution of iodide of potassium, and, lastly, some mucilage of starch; the deep-blue compound of iodine and starch is at once produced. Repeat this experiment, using nitrate of potassium instead of nitrite; no blue color is produced.

$$2HI + 2HNO_2 = 2H_2O + 2NO + I_2.$$

*Tests for Nitrites in Water.*—This liberation of iodine by nitrites and not by nitrates is a reaction of considerable value in searching for nitrites in ordinary drinking-waters, the occurrence of such salts being held to indicate the presence of nitrogenous organic matter in a state of oxidation or decay. The sulphuric acid used in the operation must be pure, and the iodide of potassium free from iodate. If much organic matter is present, however, the nitric acid liberated by the sulphuric may be reduced to nitrous acid. It is perhaps best, therefore, to add acetic acid, and (Fresenius) distil over 10 or 20 per cent. of the water and apply the test to this distillate. Very dilute solutions of nitrous acid may thus be distilled without the slightest decomposition.

*Commercial Nitrous Acid.*—The liquid commonly termed in pharmacy nitrous acid is simply nitric acid impure from the presence of nitrous acid.

*The only nitrite used in medicine* is a nitrite of an organic basylous radical, ethyl; nitrite of ethyl ($C_2H_5NO_2$), or nitrous ether, is one of the chief constituents of "*sweet spirit of nitre*" (*Spiritus Ætheris Nitrosi*, B. P. and U. S. P.: *vide* Index).

OPHELIC ACID ($C_{18}H_{20}O_{10}$).—This is one of the principles to which the herb *Ophelia chirata*, or Chiretta (*Chirata*, B. P.), owes its bitterness. It is an amorphous yellow body. Another is *Chiratin* ($C_{26}H_{46}O_{15}$), decomposable by hydrochloric acid into *Chiratogenin* ($C_{18}H_{24}O_3$) and ophelic acid (Höhn).

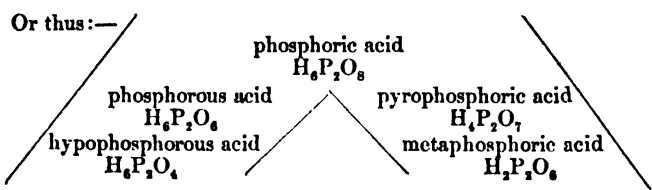
PHOSPHOROUS ACID ($H_3PO_3$, or $H_2PHO_3$).—It is necessary to notice this compound in order that the reader may have brought before him the three acids of phosphorus, namely, phosphoric acid ($H_3PO_4$), phosphorous acid ($H_2PHO_3$), and hypophosphorous acid ($HPH_2O_2$): it will be noticed that in composition they differ from each other simply in the proportion of oxygen, the molecules containing four, three, and two atoms respectively. In constitution they differ by the hypothetical phosphoric radical or grouping being trivalent, the phosphorous bivalent, and the hypophosphorous radical univalent. These three acids and corresponding salts must not be confounded with pyrophosphoric and metaphosphoric acids and salts; the former are acids of phosphorus; the latter, varieties of phosphoric acid; the former, in composition, differ from each other in the proportion of oxygen they contain; the latter, by the elements of water:—

| *Acids of Phosphorus.* | | *Varieties of Phosphoric Acid.* | |
|---|---|---|---|
| $H_3PO_4$ | phosphoric acid. | $H_3PO_4$ | (ortho)phosphoric acid. |
| $H_2PHO_3$ | phosphorous acid. | $H_4P_2O_7$ | pyrophosphoric acid. |
| $HPH_2O_2$ | hypophosphorous acid. | $HPO_3$ | metaphosphoric acid. |

When hypophosphorous acid is exposed to the air, oxygen is absorbed and phosphorous acid results; by prolonged exposure more oxygen is absorbed and phosphoric acid is obtained. When phosphoric acid, or rather, for distinction, orthophosphoric acid, is heated, every two molecules yield the elements and a molecule of water, and pyrophosphoric acid results; by prolonged exposure to heat more water is evolved, and metaphosphoric acid is obtained. These differences will be further evident if the formulæ be written empirically, nearly all being doubled, thus:—

| | |
|---|---|
| $H_6P_2O_4$ | hypophosphorous acid. |
| $H_6P_2O_6$ | phosphorous acid. |
| $H_6P_2O_8$ | phosphoric acid, or orthophosphoric acid. |
| $H_4P_2O_7$ | pyrophosphoric acid. |
| $H_2P_2O_6$ | metaphosphoric acid. |

Or thus:—



From the central compound, phosphoric acid, the acids of phosphorus differ by regularly diminishing proportions of the element oxygen (see previous page), the varieties of phosphoric acid by regularly diminishing proportions of the elements of water.

*Prepare phosphorous acid* by exposing a moist stick of phosphorus to the air; a thin stream of heavy white vapor falls, which contains the acid in question. The best method of collection is to place the stick in an old test-tube having a hole in the bottom, to support this tube by a funnel or otherwise, the neck of the funnel being supported in a bottle, test glass, or tube, at the bottom of which is a little water. Having collected some phosphorous acid in this way, apply the various tests already alluded to under *Hypophosphorous Acid*, first carefully neutralizing the phosphorous acid by an alkali. The means by which the varieties of phosphoric acid are distinguished have been given under *Metaphosphoric Acid.* Associated with the phosphorous acid, prepared as above stated, there is said to be an acid having the formula $H_2PO_3$, and termed *hypophosphoric acid.* Its anhydride would be $P_2O_4$.

Other soluble phosphites are prepared by neutralizing phosphorous acid with alkalies, and the insoluble phosphites by double decomposition.

PYROGALLIC ACID.—See *Tannic Acid.*

PYROPHOSPHORIC ACID ($H_4P_2O_7$) AND OTHER PYROPHOSPHATES.—Heat ordinary phosphate of sodium ($Na_2HPO_4$, $12H_2O$) in a crucible; water of crystallization is first evolved, and dry phosphate ($Na_2HPO_4$) remains. Continue the heat to redness; two molecules of the salt yield one molecule of water, and a salt having new properties is obtained:—

$$2Na_2HPO_4 - H_2O = Na_4P_2O_7.$$

It is termed pyrophosphate of sodium, in allusion to its origin (πῦρ, *pūr*, fire). Phosphoric acid itself is similarly affected by heat, $2H_3PO_4 - H_2O = H_4P_2O_7$ (pyrophosphoric acid), though metaphosphoric acid is also formed. Other pyrophosphates are produced in a similar way, or by double decomposition and precipitation, or by neutralizing pyrophosphoric acid by an oxide, hydrate, or carbonate. Possibly the pyrophosphates are only compounds of orthophosphates with metaphosphates:—

$$Na_4P_2O_7 = Na_3PO_4, NaPO_3.$$

*Tests.*—To solution of a pyrophosphate add solution of nitrate of silver; white pyrophosphate of silver ($Ag_4P_2O_7$) falls as a dense white powder, differing much in appearance from the white gelatinous metaphosphate of silver or the yellow ortho-

phosphate. To pyrophosphoric acid or to a pyrophosphate mixed with acetic acid, add an aqueous solution of albumen (white of egg); no precipitate occurs. Metaphosphoric acid, it will be remembered, gives a white precipitate with albumen.

## QUESTIONS AND EXERCISES.

600. What are the sources of lactic acid?
601. How is lactic acid usually prepared?
602. Name some of the plants in which malic acid is found.
603. Whence is meconic acid derived?
604. By what process may meconic acid be isolated?
605. Which is the best test for the meconic radical?
606. Distinguish meconates from sulphocyanates.
607. Give the mode of manufacture of hypophosphites.
608. How is phosphoretted hydrogen prepared?
609. By what ready method may metaphosphoric acid be obtained for experimental purposes?
610. Name the tests for metaphosphates.
611. How may meta- or pyro-phosphoric acid be converted into orthophosphoric acid?
612. Describe the preparation of phosphorous acid.
613. State the relations which the acids of phosphorus bear to each other.
614. How are pyrophosphates prepared?
615. Offer two views of the constitution of pyrophosphates.
616. Define, by formulæ, metaphosphates, pyrophosphates, orthophosphates, phosphites, and hypophosphites.
617. Mention the tests by which meta-, pyro-, and orthophosphates are analytically distinguished.
618. Name the reactions by which hypophosphites and phosphites are detected.

SILICIC ACID ($H_4SiO_4$) AND OTHER SILICATES.—Silicates of various kinds are among the commonest of minerals. The various *clays* are aluminium silicates; the volcanic substance termed *pumice-stone* is a porous silicate of aluminium and of alkali-metals or alkaline-earth metals; *meerschaum* is an acid silicate of magnesium; the ordinary *sandstones* are chiefly silica; *sand*, *flint*, *quartz*, *agate*, *chalcedony*, and *opal* are silicic anhydride or *silica* ($SiO_2$). *Tripoli*, a polishing powder now found in many other countries than Tripoli, consists of infusorial skeletons of nearly pure silica. *Bath brick*, used in knife-polishing, is a silico-calcareous deposit found in the estuary at Bridgewater and other places. *Asbestos* or *amianth* is a fibrous silicate of calcium and magnesium, the length of the fibres being from less than one inch to five feet. A single silk-like fibre can easily be fused, but, even in very small masses, and for all practical

purposes, asbestos is infusible and, of course, incombustible. It is also a bad conductor of heat. It is already largely used in piston-rods and joints and for steam apparatus generally; as a covering for boilers to prevent loss of heat by radiation; and for so lining ceilings, floors, and other partitions as to render rooms, etc., fireproof. Artificial silicates are familiar under the forms of *glass* and *earthenware*. Common English window glass is usually silicate of calcium, sodium, and aluminium; French glass, silicate of calcium and sodium; Bohemian, chiefly silicate of potassium and calcium; English flint- or crystal-glass for ornamental, table, and optical purposes, is mainly silicate of potassium and lead. Earthenware is mostly silicate of aluminum (clay), with more or less of silicate of calcium, sodium and potassium, and, in the commoner forms, silicates of iron. The various kinds of *porcelain* (China, Sèvres, Meissen, Berlin, English), *Wedgwoodware*, and *stoneware* are varieties of earthenware. *Crucibles*, *bricks*, and *tiles* are clay silicates. *Mortar* is essentially silicate of calcium. *Portland*, *Roman*, and other hydraulic *cements* are silicates of calcium with more or less silicate of aluminium.

Mix together a few grains of powdered flint or sand with about five or six times its weight of carbonate of sodium and an equal quantity of carbonate of potassium, and fuse a little of the mixture on platinum-foil in the blowpipe-flame; the product is a kind of *soluble glass*. Boil the foil in water for a few minutes, filter; to a portion add excess of hydrochloric acid, evaporate the solution to dryness, and again boil the residue in water and acid; oxide of silicon, *silicic anhydride*, or *silica* ($SiO_2$), remains as a light, flaky, insoluble powder.

The *soluble glass* or *glass liquor* of trade commonly contains 10 or 12 per cent. of soda (NaHO) to 20 or 25 per cent. of silica ($SiO_2$).

The foregoing operation constitutes the *test* for silicates. By fusion with alkali the silicate is decomposed, and a soluble alkaline silicate formed. On addition of acid, silicic acid ($H_4SiO_4$) is set free, but remains in solution if sufficient water is present. The heat subsequently applied eliminates water and reduces the silicic acid to silica ($SiO_2$), which is insoluble in water or acid. By the addition of hydrochloric acid to soluble glass, and removal of the resulting alkaline chloride and excess of hydrochloric acid by dialysis (a process to be subsequently described), a pure aqueous solution of silicic acid may be obtained; it readily changes into a gelatinous mass of silicic acid. Possibly some of the natural crystallized varieties of silica may have been obtained from the silica contained in such an aqueous solution, nearly all waters yielding a small quantity of silica when treated as above described.

*A variety of silicic acid* ($H_2SiO_3$) sometimes termed *dibasic*, to distinguish it from the normal or *tetrabasic* acid ($H_4SiO_4$), results when the aqueous solution of the latter is evaporated *in vacuo*.

*Siliciuretted hydrogen*, or hydride of silicon ($SiH_4$), is a spontaneously inflammable gas formed on treating silicide of magnesium

with hydrochloric acid. It is the analogue of light carburetted hydrogen ($CH_4$). A liquid chloride of silicon ($SiCl_4$) and a gaseous fluoride ($SiF_4$) also exist. Many other analogies are traceable between the elements silicon, boron, and carbon (see p. 333).

Succinic Acid ($H_2C_4H_4O_4$).—Amber (*Succinum*) is a peculiar resin usually occurring in association with coal and lignite. From the fact that fragments of coniferous fruit are frequently found in amber, and impressions of bark on its surface, it is considered to have been an exudation from a species of Pinus now probably extinct. Heated in a retort, amber yields, first, a sour aqueous liquid containing acetic acid and another characteristic body appropriately termed *succinic acid;* second, a volatile liquid known as *oil of amber* (*Oleum Succini Rectificatum*, U. S. P.), resembling the oil yielded by most resinous substances under similar circumstances; and, third, a pitchy residue allied to asphalt. The succinic acid is a normal constituent of the amber, the acetic acid is produced during distillation. Succinic acid has also been found in wormwood, in several pine-resins, and in certain animal fluids, such as those of hydatid cysts, and hydrocele. It may be obtained artificially from butyric, stearic, or margaric acid by oxygen. Tartaric, malic, and succinic acids are also convertible the one into the other.

The *succinates* are normal ($R'_2C_4H_4O_4$) and acid ($R'HC_4H_4O_4$); a double succinate of potassium and hydrogen ($KHC_4H_4O_4.H_2C_4H_4O_4.H_2O$), analogous to the superacid oxalate, salt of sorrel, also exists.

Soluble succinates give a bulky brown precipitate with neutral ferric chloride, only less voluminous than ferric benzoate; a white precipitate with acetate of lead, soluble in excess of either reagent; with nitrate of silver a white precipitate after a time; with chloride of barium no precipitate at first, but a white one of succinate of barium on the addition of ammonia and alcohol. Succinates are distinguished from benzoates by the last-named reaction, and by not yielding a precipitate on the addition of acids (*vide* p. 336).

Sulphocyanic Acid (HCyS) and other Sulphocyanates.—Boil together sulphur and solution of cyanide of potassium; solution of sulphocyanate of potassium (KCyS) is formed. Warm the liquid, add hydrochloric acid till it faintly reddens litmus paper, and filter; any sulphide of potassium is thus decomposed, and the solutions may then be used for the following reactions.

*Tests.*—Filter, and to a small portion of the solution add a ferric salt ($Fe_2Cl_6$); a deep blood-red solution of ferric sulphocyanate is formed. To a portion of the red liquid add a little hydrochloric acid; the color is not discharged (meconate of iron, a salt of similar tint, is decomposed by hydrochloric acid). In the acid liquid place a fragment or two of zinc; sulphuretted hydrogen is evolved, and the red color disappears. To another

portion of the ferric sulphocyanate add solution of corrosive sublimate; the color is at once discharged. (Ferric meconate is unaffected by corrosive sublimate.) The ferric is the best test of the presence of a sulphocyanate; indirectly, it is a good test of the presence of hydrocyanic acid or cyanogen. Solutions of pure ferrous salts are not colored by the solution of sulphocyanate.

To solution of a sulphocyanate add solution of mercuric nitrate; mercuric sulphocyanate is precipitated as a white powder.

*Pharaoh's Serpents.*—Mercuric sulphocyanate, thoroughly washed and made up into little cones, forms the toy called Pharaoh's serpent. It readily burns when ignited, the chief product being a light solid matter (mellon, $C_9N_{13}$, and the melam, $C_3H_6N_6$), which issues from the cone in a snake-like coil of extraordinary length. The other products are mercuric sulphide (of which part remains in the snake and part is volatilized), nitrogen, sulphurous, and carbonic acid gases, and vapor of metallic mercury. (For details concerning the economical manufacture of sulphocyanates, see *Pharmaceutical Journal*, second series, vol. vii. p. 581, and p. 152.)

The sulphocyanic radical (CyS) is often termed sulphocyanogen (Scy), and its compounds regarded as *sulphocyanides*. Saliva contains sulphocyanides.

TANNIC ACID, or TANNIN ($C_{27}H_{22}O_{17}$ or perhaps $C_{14}H_{17}O_9$).—This is a common astringent constituent of plants, but is contained in largest quantity in galls (excrescences on the oak formed by the puncture and deposited ova of an insect). English galls contain from 14 to 28 per cent. of tannic acid; Aleppo galls (*Galla*, B. P. and U. S. P.) from 25 to 65 per cent. (*Acidum Tannicum*, B. P. and U. S. P.)

*Process.*—"Expose powdered galls (about an ounce is sufficient for the purpose of study) to a damp atmosphere for two or three days, and afterwards add sufficient ether to form a soft paste. Let this stand in a well-closed vessel for twenty-four hours, then having quickly enveloped it in a linen cloth, submit it to strong pressure so as to separate the liquid portion, which contains the bulk of the tannin in solution. Reduce the pressed cake to powder, mix it with sufficient ether, to which one-sixteenth of its bulk of water has been added, to form again a soft paste, and press this as before. Mix the expressed liquids, and expose the mixture to spontaneous evaporation until, by the aid subsequently of a little heat, it has acquired the consistence of a soft extract; then place it on earthen plates or dishes, and dry it in a hot-air chamber at a temperature not exceeding 212°."

The resulting tannic acid occurs in "pale yellow vesicular

masses or thin glistening scales, with a strongly astringent taste, and an acid reaction, readily soluble in water and rectified spirit, very sparingly soluble in pure ether," though soluble in the ethereal fluid used in the foregoing process—a fluid which is really a mixture of true ether, water, and alcohol (both the latter contained in the common "ether"), and a little added water also.

*Medicinal Uses.*—Tannic acid is very soluble in water, and in this form is usually administered in medicine. Its official preparations are *Glyceritum Acidi Tannici*, *Suppositoria Acidi Tannici*, and *Trochisci Acidi Tannici*.

*Tests.*—To an aqueous solution of tannic acid add aqueous solution of gelatine, a yellowish-white flocculent compound of the two substances is precipitated. This is a good test of the presence of tannic acid.

*Tanning.*—The above reaction also serves to explain the chemical principle involved in *tanning*—the operation of converting skin into leather. In that process the skin is soaked in infusion of oak-bark (*Quercus Cortex*), the tannic acid of which uniting with the gelatinous tissues of the skin yields a compound very well represented by the above precipitate. The outer bark of the oak contains little or no tannic acid, and is commonly shaved off from the pieces of bark which are large enough to handle; useless coloring matter is thus also rejected. Other infusions and extracts besides that of oak-bark (chiefly catechu, sumach, and valonia) are largely used by tanners; if used alone these act too quickly, and give a harsh, hard, less durable leather. The tannic acid of these preparations is probably slightly different from that of oak-bark.

To an aqueous solution of tannic acid add a neutral solution of a ferric salt; dark bluish-black tannate of iron is slowly precipitated. This is an excellent test for the presence of tannic acid in vegetable infusions. The precipitate is the basis of nearly all black writing-inks. Ferrous salts give at first only a slight reaction with tannic acid; but the liquid gradually darkens. Characters written with this liquid become quite black in a few hours, and are very permanent.

To an aqueous solution of tannic acid add solution of tartar-emetic; tannate of antimony is precipitated. This reaction and that with gelatine are useful in the quantitative estimation of the amount of tannic acid in various substances.

*Tannic acid* as it occurs in oak-bark is said to be a glucoside, that is, like several other substances, yields glucose (grape-sugar) when boiled with dilute sulphuric or hydrochloric acid, the other product being gallic acid:—

$$C_{27}H_{22}O_{17} + 4H_2O = C_6H_{12}O_6 + 3H_3C_7H_3O_5.$$

*Catechu* (*Catechu pallidum*, B. P.), *Gambier*, or *Terra Japonica*, an extract of the *Uncaria Gambier;* as well as the true *Catechu*, *Cutch*, or *Terra Japonica*, an extract from the *Acacia Catechu* (*Catechu nigrum*, P. I.) and *A. Suma; East Indian Kino* (*Kino*, B. P.) from the *Pterocarpus marsupium;* also *Bengal* or *Butea Kino*, from the *Palas* or *Dhak* tree, *Butea frondosa* (*Buteæ gummi vel Kino Bengalensis*, P. I.); *Elm-bark* (*Ulmi Cortex*, B. P.); and some other vegetable products, contain a variety of tannic acid (*mimotannic acid*), which gives a greenish precipitate with neutral solutions of ferric salts. According to Paul and Kingzett it yields, when decomposed, unfermentable sugar, and an acid different from ordinary gallic acid. Catechu and Gambier also contain *catechuic acid* or *catechin*, $C_{15}H_{12}O_5$, a body occurring in minute colorless acicular crystals, and, like mimotannic acid, affording a green precipitate with ferric salts.

*Bael fruit* (*Belæ Fructus*, B. P.), from the *Ægle Marmelos*, is said to owe its power as a remedy for dysentery and diarrhœa to a variety of tannic acid, but this is questionable. The rind of the fruit of the *pomegranate* (*Punica granatum*) (*Granati Cortex*, P. I.) contains tannic acid. The astringency of Pomegranate-root Bark (*Granati Radicis Cortex*, B. P. and P. I.) is due to a tannic acid (its anthelmintic properties probably to a resinoid matter or possibly to what Tanret states to be an alkaloid *pelletierine*). A tannic acid also probably gives the astringency to Logwood (*Hæmatoxyli Lignum*, B. P., the *color* of which is due to oxidized *hæmatoxylin*). *Rhatany*-root bark (*Krameriæ Radix*, B. P.) contains about 20 per cent. of tannic acid, its active astringent principle; rhubarb root about 9 per cent. *Bearberry*-leaves (*Uvæ Ursi Folia*, B. P.) owe most of their therapeutic power to about 35 per cent. of tannic acid. (The cause of their influence on the kidneys is not yet traced.) They also contain *arbutin*, a crystalline glucoside. *Larch bark* (*Laricis Cortex*, B. P.), the inner bark of *Pinus larix* or *Larix europæa*, contains, according to Stenhouse, a considerable amount of a tannic acid giving olive-green precipitates with salts of iron, and *larixin* and *larixinic acid* ($C_{10}H_{10}O_5$), a somewhat bitter substance. *Areca nuts* or *Betel nuts* (*Areca*, B. P.), from the Areca Palm (*Areca catechu*), contain, according to Flückiger and Hanbury, about 15 per cent. of "tannic matter." The extract of the fruit of *Gab*, or *Diospyros embryopteris* (*Diospyri Fructus*, P. I.), is a powerful astringent containing tannic acid. The rhizome of *Geranium maculatum*, Spotted Cranesbill or Alum-root, contains both tannic and gallic acids.

*Gallic Acid* ($H_3C_7H_3O_5,H_2O$) (*Acidum Gallicum*, B. P. and U. S. P.) occurs in small quantity in oak-galls and other vegetable substances, but is always prepared from tannic acid. Powdered galls are moistened with water and set aside in a warm place for five or six weeks, occasionally being remoistened; fermentation occurs, and impure gallic acid is formed. The product is treated with about three times its weight of water, boiled to dissolve the gallic acid, filtered, the solution set aside

to cool, deposited gallic acid collected, drained, pressed between folds of paper to remove all mother-liquor, and, if necessary, purified by recrystallization from water, or by solution in hot water with animal charcoal, which absorbs coloring matter. On filtering and cooling, most of the acid separates in the form of fawn-colored slender acicular crystals. Gallic acid is soluble in about 100 times its weight of cold or 3 of boiling water, freely in spirit, sparingly in ether, also in glycerine (*Glyceritum Acidi Gallici*, U. S. P.).

The nature of the action by which gallic acid is thus produced is probably similar to that of the action of dilute acids on tannic acid. During the process oxygen is absorbed and carbonic acid gas evolved, the sugar being thus broken up or perhaps prevented from being formed.

*Test.*—To an aqueous solution of gallic acid add a neutral solution of ferric salt; a bluish-black precipitate of gallate of iron falls, similar in appearance to tannate of iron. Ferrous salts are also blackened by gallic acid. To more of the solution add an aqueous solution of gelatine; no precipitate occurs. By the latter test gallic acid is distinguished from tannic acid.

*Pyrogallic Acid* ($C_6H_6O_3$).—This substance sublimes in light feathery crystals when gallic acid is heated. To an aqueous solution add a neutral solution of a ferric salt, a red color is produced. To another portion add a ferrous salt; a deep-blue color results.

*Test for the three acids.*—To three separate small quantities of milk of lime in test-tubes add, respectively, tannic, gallic, and pyrogallic acids; the first slowly turns brown, the second more rapidly, while the pyrogallic mixture at once assumes a beautiful purplish-red color changing to brown. These reactions are highly characteristic. They are accompanied by absorption of oxygen from the air.

*Use of Pyrogallic Acid in Gas-analysis.*—A mixture of pyrogallic acid and solution of potash absorbs oxygen with such rapidity and completeness that a strong solution of each, passed up successively by a pipette into a graduated tube containing air or other gas, forms an excellent means of estimating free oxygen. The value of this method may be roughly proved by pouring a small quantity of each solution into a phial, immediately and firmly closing its mouth with a cork, thoroughly shaking the mixture and then removing the cork under water; the water rushes in and occupies about one-fifth of the previous volume of air, indicating that the atmosphere contains one-fifth of its bulk of oxygen. The small amount of carbonic acid gas present in the air is also absorbed by the alkaline liquid; in delicate experiments this should be removed by the alkali before the addition of pyrogallic acid.

URIC ACID ($H_3C_5H_2N_4O_3$) AND OTHER URATES.—Acidulate a few ounces of human urine with hydrochloric acid, and set aside for twenty-four hours; a few minute crystals of uric acid

will be found adhering to the sides and bottom of the vessel and floating on the surface of the liquid.

*Microscopical Test.*—Remove some of the floating particles by a slip of glass, and examine by a powerful lens or microscope; the chief portion will be found to be in yellowish semitransparent crystals, more or less square, two of the sides of which are even, and two very jagged; but other forms are common (see the lithographs in the section on Urinary Sediments).

*Chemical Test.*—Collect more of the deposit, place in a watch-glass or small white evaporating-dish, remove adherent moisture by a piece of blotting- or filter-paper, add a drop or two of strong nitric acid, and evaporate to dryness; the residue will be red. When the dish is cold, add a drop of solution of ammonia; a purplish-crimson color results. The color is deepened on the addition of a drop of solution of potash.

*Notes.*—Uric acid (or lithic acid) and urates (or lithates) of sodium, potassium, calcium, and ammonium, are common constituents of animal excretions. Human urine contains about one part of urate (usually urate of sodium) in 1000. When more than this is present the urate is often deposited as a sediment in the excreted urine, either at once, or after standing a short time. Uric acid or other urate is also occasionally deposited before leaving the bladder, and, slowly accumulating there, forms a common variety of urinary calculus.——Some urates are not definitely crystalline; but, when treated with dilute nitric acid or a drop of solution of potash, and then a drop or two of acetic acid, jagged microscopic crystals of uric acid are usually formed.—All urates yield the crimson color when treated as above described.—This color is due to a definite substance *murexid* ($C_8H_8N_6O_6$) (from the *murex*, a shell-fish of similar tint); and the test is known as *the murexid test.* The formation of murexid is due to the action of ammonia on *alloxan* ($C_4H_2N_2O_4,4H_2O$) and other white crystalline products of the oxidation of uric acid by nitric acid. Murexid is a good dye; it may be prepared from *guano* (the excrement of sea-fowl), which contains a large quantity of urate of ammonium. The excrement of the serpent is almost pure ammonium urate.

Uric acid and the urates will be again alluded to in connection with the subject of morbid urine.

VALERIANIC ACID OR VALERIC ACID ($HC_5H_9O_2$) AND OTHER VALERIANATES.—In a test-tube place a few drops of amylic alcohol (fousel oil) with a little dilute sulphuric acid and a grain or two of red chromate of potassium, cork the tube, set aside for a few hours, and then heat the mixture; valerianic acid, of characteristic valerian-like odor, is evolved.

Valerianic acid occurs naturally in valerian-root in association with the essential oil from which it is derived (*vide* Index), but is usually prepared artificially, by the foregoing process, from amylic

alcohol, to which it bears the same relation as acetic acid does to common alcohol:—

$$C_2H_5HO + O_2 = HC_2H_3O_2 + H_2O$$
$$C_5H_{11}HO + O_2 = HC_5H_9O_2 + H_2O.$$

*Valerianate of Sodium* ($NaC_5H_9O_2$) (*Sodæ Valerianas*, B.P.) is prepared from the valerianic acid and valerianate of amyl obtained on distilling the mixture of amylic alcohol (4 fl. ozs.), sulphuric acid (6½ fl. ozs. with 10 of water), and red chromate of potassium (9 ozs. in 70 of water). The mixture should stand for several hours before heat is applied.

$$\underset{\text{Red chromate of potassium.}}{2(K_2CrO_4,CrO_3)} + \underset{\text{Sulphuric acid.}}{8H_2SO_4} = \underset{\text{Sulphate of potassium and chromium. (Chrome alum.)}}{2(K_2SO_4,Cr_2SO_4)} + \underset{\text{Water.}}{8H_2O} + \underset{\text{Oxygen.}}{3O_2}$$

$$\underset{\text{Amylic alcohol.}}{C_5H_{11}HO} + \underset{\text{Oxygen.}}{O_2} = \underset{\text{Valerianic acid.}}{HC_5H_9O_2} + \underset{\text{Water.}}{H_2O}$$

$$\underset{\text{Amylic alcohol.}}{2C_5H_{11}HO} + \underset{\text{Oxygen.}}{O_2} = \underset{\text{Valerianate of amyl.}}{C_5H_{11}C_5H_9O_2} + \underset{\text{Water.}}{2H_2O}$$

The distillate (70 or 80 ozs.) is saturated with soda, which not only yields valerianate of sodium with the free valerianic acid, but decomposes the valerianate of amyl produced at the same time, more valerianate of sodium being formed and some amylic alcohol set free, according to the following equations:—

$$\underset{\text{Valerianic acid.}}{HC_5H_9O_2} + \underset{\text{Soda.}}{NaHO} = \underset{\text{Valerianate of sodium.}}{NaC_5H_9O_2} + \underset{\text{Water.}}{H_2O}$$

$$\underset{\text{Valerianate of amyl.}}{C_5H_{11}C_5H_9O_2} + \underset{\text{Soda.}}{NaHO} = \underset{\text{Valerianate of sodium.}}{NaC_5H_9O_2} + \underset{\text{Amylic alcohol.}}{C_5H_{11}HO}$$

From the solution of valerianate of sodium (which should be made neutral to test-paper by careful addition of soda solution) the solid white salt is obtained by evaporation to dryness and cautious fusion of the residue. The mass obtained on cooling should be broken up and kept in a well-closed bottle. It is entirely soluble in spirit.

*Other Valerianates*, as valerianate of zinc (*Zinci Valerianas*, B. P. and U. S. P.), and ferric valerianate, may be made by double decomposition of valerianate of sodium with the sulphate or other salt of the metal the valerianate of which is desired, the new valerianate either precipitating or crystallizing out. A hot solution of sulphate of zinc (5¾ parts) and valerianate of sodium (5 parts) in water (40 parts) gives a crop of crystals of valerianate of zinc on cooling.

*Tests.*—Heated with diluted sulphuric acid, valerianates of the metals give valerianic acid, which has a highly character-

istic smell. Valerianate of sodium thus treated, and the resulting oily acid liquid purified by agitation with sulphuric acid and distillation, furnishes the *Acidum Valerianicum* of the United States Pharmacopœia (sp. gr. .933). Dry ammonia gas passed into valerianic acid gives white lamellar crystals of valerianate of ammonium (*Ammonii Valerianas*, U. S. P.).

The amylic alcohol ($C_5H_{11}HO$) from which valerianates are prepared may contain the next lower homologue, *butylic alcohol* ($C_4H_9HO$). This, during oxidation, will be converted into *butyric acid* ($HC_4H_7O_2$), the next lower homologue of valerianic acid ($HC_5H_9O_2$), and hence the various valerianates be contaminated by some *butyrates*. These are detected by distillation with diluted sulphuric acid and addition of solution of acetate of copper to the distillate, which at once becomes turbid if butyric acid be present. In this reaction valerianic and butyric acids are produced by double decomposition of the valerianate and butyrate by the sulphuric acid, and distil over on the application of heat. On the addition of acetate of copper ($Cu2C_2H_3O_2$) butyrate of copper ($Cu2C_4H_7O_2, H_2O$) is formed, and, being almost insoluble in water, is at once precipitated, or remains suspended, giving a bluish-white opalescent liquid. Valerianate of copper ($Cu2C_5H_9O_2$) is also formed after some time, but is far more soluble than the butyrate, and only slowly collects in the form of greenish oily drops, which gradually pass into greenish-blue hydrous crystalline valerianate of copper (Larocque and Huralt).

VANILLIC ACID ($HC_8H_7O_3$) or VANILLIN ($C_8H_8O_3$) or METHYL-PROTOCATECHUIC ALDEHYDE ($C_7H_5CH_3O_3$); the body to which is due the odor and flavor of vanilla.—The white crystals commonly found on vanilla—the prepared unripe pods of *Vanilla planifolia*—previously termed *vanillin*, were found by Carles to be a weak acid. It occurs in vanilla to the extent of from 1½ to 3 per cent. Vanillin has recently been prepared artificially by Tiemann and Haarmann from coniferin, a glucoside existing in the sapwood of pines. The body remaining after the removal of glucose from coniferin, or, indeed, coniferin itself, by action of a mixture of red chromate of potassium and sulphuric acid, yields the vanillin. It also may be obtained by a series of reactions starting from that of carbonic acid on carbolate of potassium; also from the eugenol of oil of cloves. By action of hydrochloric acid, vanillin yields chloride of methyl and pyrocatechuic aldehyd. Such reactions will be better understood when the pupil has studied succeeding sections on what is commonly termed *Organic Chemistry*. Artificial vanillin is less stable than natural vanillin, perhaps because with the latter is associated a preservative resin.

## QUESTIONS AND EXERCISES.

619. What is the constitution of nitrites?
620. Mention a test for nitrites in potable waters.
621. Which nitrite is official?
622. Give the names of some natural and artificial silicates.
623. What is "soluble glass"?
624. Distinguish between silica and silicic acid.
625. How are silicates detected?
626. What is the quantivalence of silicon?
627. Mention the sources, formulæ, and analytical reactions of succinates.
628. State the mode of manufacture and tests of sulphocyanates.
629. What proportion of tannic acid is contained in galls?
630. Describe the official process for the preparation of Tannic Acid.
631. Explain the chemistry of "tanning."
632. Enumerate the tests for tannic acid.
633. What is the assumed constitution of tannic acid?
634. Mention other official substances whose astringency is due to tannic acid.
635. How is gallic acid prepared?
636. By what reaction is gallic distinguished from tannic acid?
637. Mention the characteristic properties of pyrogallic acid.
638. Explain the murexid test for uric acid.
639. Describe the artificial preparation of valerianic acid and other valerianates, giving diagrams or equations.
640. What is the formula of valerianic acid?
641. How are butyrates detected in presence of valerianates?

---

## DETECTION OF THE ACIDULOUS RADICALS OF SALTS SOLUBLE IN WATER.

Analytical operations may now be resumed, the detection of acidulous radicals being practised for two or three days, and then full analyses made, both for basylous and acidulous radicals. To this end a few compounds of stated metals (potassium, sodium, or ammonium) should be placed in the hands of the practical student for examination according to the following paragraphs and Tables. Mixtures in which both basylous and acidulous radicals may be sought should then be analyzed.

In examining salts soluble in water, and concerning which no general information is obtainable, search must first be made for any basylous radicals by the appropriate methods (*vide* page 221 or 258). Certain metals having been thus detected, a little reflection on the character of their salts will at once indicate what acidulous radicals may be, and what cannot be, present. Thus, for instance, if the sub-

stance under examination is freely soluble in water, and lead is found, only the nitric and acetic radicals need be sought, none other of the lead salts than nitrate or acetate being freely soluble in water. Moreover, the salt is more likely to be acetate than nitrate of lead, for two reasons: the former is more soluble than the latter, and is by far the commoner salt of the two. Medical and pharmaceutical students have probably, in dispensing, already learned much concerning the solubility of salts, and whether a salt is rarely employed or in common use. And although but little dependence can be placed on the chances of a salt being present or absent according to its rarity, still the point may have its proper weight. If, in a mixture of salts, ammonium, potassium, and magnesium have been found associated with the sulphuric, nitric, and hydrochloric radicals, and we are asked how we suppose these bodies may exist in the mixture, it is far more in accordance with common sense to suggest that sal-ammoniac, nitre, and Epsom salt were originally mixed together than to suppose any other possible combination. Such appeals to experience regarding the solubility or rarity of salts cannot be made by any one not previously acquainted, or insufficiently acquainted, with the characters of salts; in such cases the relation of a salt to water and acids can be ascertained by referring to the following Table (p. 365) of the solubility or insolubility of about five hundred of the common or rarer salts met with in chemical operations.

The opposite course to the above (namely, to ascertain what acidulous radicals are present in a mixture, and then to appeal to experience to tell what basylous radicals may be and what cannot be present) is impracticable; for acidulous radicals cannot be separated out, one after the other, from one and the same quantity of substance by a similar treatment to that already given for basylous radicals. Indeed such a sifting of acidulous radicals could scarcely be accomplished at all, or only by a vast deal of labor. The basylous radicals must, therefore, be first detected.

Even when the basylous radicals have been found, the acidulous radicals which may be present must be sought for singly, the only additional aid which can be brought in being the action of sulphuric acid, a barium salt, a calcium salt, nitrate of silver, and ferric chloride on *separate* small portions of the solution under examination, as detailed in the second of the following Tables.

*Commence the analysis* of an aqueous solution of a salt or salts, the basylous radicals in which are known, by writing out a list of the acidulous radicals which may be, or, if more convenient, of those which cannot be present. To this end consult the following Table (p. 365) of the solubility of salts in water. Look for the name of the metal of the salt in the vertical column; the letters S and I indicate which salts are soluble and which insoluble in water, an asterisk attached to the S meaning that the salt is slightly soluble. The acidulous part of the name is given in the top line of the Table. All the names are in alphabetical order, for facility of reference.

Some of the salts marked as insoluble in water are soluble in aqueous solutions of soluble salts, a few forming soluble double salts. To characterize salts as soluble, slightly soluble, or insoluble, only roughly indicates their relation to water: on the one hand, very few salts are absolutely insoluble in water; on the other there is a limit to the solubility of every salt.

*If only one, two, or perhaps three given acidulous radicals can be in the liquid, test directly for it or them according to the reactions given in the previous pages. If several may be present, pour small portions of the solutions, rendered neutral if necessary by ammonia, into five test-tubes, and add respectively sulphuric acid, nitrate or chloride of barium, chloride of calcium, nitrate of silver, and ferric chloride; then consult the Table on page* 366, *in order to correctly interpret the effects these reagents may have produced.*

TABLE OF THE SOLUBILITY OR INSOLUBILITY OF SALTS IN WATER.

| | Acetate. | Arsenate. | Arsenite. | Carbonate. | Chloride. | Citrate. | Chromate. | Cyanide. | Hydrate. | Iodide. | Nitrate. | Oxalate. | Oxide. | Phosphate. | Sulphate. | Sulphide. | Sulphite. | Tartrate. |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| Aluminium | S | I | I | I | S | S* | I | - | I | - | S | I | I | I | S | I | I | S |
| Ammonium | S | S | S | S | S | - | S | S | S | S | S | S | - | S | S | S | S | S |
| Antimony | S | I | I | - | S | - | I | - | I | I | S | I | S | I | S | I | I | I |
| Barium | S | I | I | - | S | S* | I | S | S | I | S | I | I | I | S | I | I | I |
| Bismuth | S | I | - | I | S | S* | I | - | I | I | S | I | I | I | S | I | I | I |
| Cadmium | S | I | - | I | S | S* | I | - | I | S | S | I | I | I | S | S | I | S* |
| Calcium | S | I | I | I | S | S* | S | S | I | S | S | I | I | I | S | I | I | S* |
| Chromium | S | I | I | I | S | I | I | I | I | S | S | I | I | I | S | I | I | S |
| Cobalt | S | I | I | I | S | S | I | I | I | S | S | I | I | I | S | I | I | S |
| Copper | S | I | I | I | S | S | I | I | I | S | S | I | I | I | S | I | I | S* |
| Ferric | S | I | I | I | S | S | S | I | I | S | S | S* | I | I | S | I | I | S |
| Ferrous | S | I | I | I | S | S | I | I | I | S | S | S* | I | I | S | I | I | S* |
| Gold | S | I | I | I | S | - | I | I | I | I | S | I | I | I | S | I | I | - |
| Lead | S | I | I | I | S* | S* | I | I | S* | S* | S | I | I | I | S | I | I | I |
| Magnesium | S | I | I | I | S | S | S | I | S | S | S | I | I | I | S | S | S | S |
| Manganese | S | I | I | I | S | I | S | I | I | S | S | I | I | I | S | I | I | S* |
| Mercuric | S | I | I | I | S | I | S* | I | I | I | S | I | I | I | S | I | I | I |
| Mercurous | S* | I | I | I | I | I | I | I | I | I | S | I | I | I | S* | I | I | I |
| Nickel | S | I | I | I | S | S | I | I | I | S* | S | I | I | I | S | I | I | S* |
| Platinum | - | I | I | - | S | S | I | - | I | S* | S | I | I | - | S | S | S | - |
| Potassium | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S |
| Silver | S* | I | I | I | I | I | I | I | S | I | S | I | I | I | S | I | I | I |
| Sodium | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S |
| Stannic | S | I | I | - | S | - | I | - | I | S | S | S | I | I | S | I | I | - |
| Stannous | S | I | I | - | S | - | I | - | I | S | S | S | I | I | S | I | I | S |
| Strontium | S | I | I | I | S | I | I | I | I | S | S | I | I | I | S | I | I | I |
| Zinc | S | I | - | I | S | S* | I | I | - | S | S | I | I | I | S | I | I | S* |

## TABLE TO AID IN THE DETECTION OF CHLORIDES, BROMIDES, IODIDES, CYANIDES, NITRATES, CHLORATES, BORATES, ACETATES, SULPHIDES, SULPHITES, SULPHATES, CARBONATES, OXALATES, TARTRATES, PHOSPHATES, AND CITRATES, IN A NEUTRAL AQUEOUS SOLUTION.

(For remarks concerning this Table see next page.)

| Sulphuric acid decomposes | Chloride of Barium precipitates | Chloride of Calcium precipitates | Nitrate of Silver precipitates | Ferric Chloride precipitates | Not precipitated |
|---|---|---|---|---|---|
| Sulphides.<br>Sulphites.<br>Carbonates, with effervescence—hydrosulphuric, sulphurous, and carbonic acid gases, known by smell, being evolved.<br>Cyanides, with production of the odor of hydrocyanic acid.<br>Acetates, with production of the odor of acetic acid when the solution is warmed. | Borates.<br>Sulphites.<br>Sulphates.<br>Carbonates.<br>Oxalates.<br>Tartrates.<br>Phosphates.<br>Citrates.<br>Of these white barium precipitates, the sulphate is the only one insoluble in hydrochloric acid; the tartrate and citrate char when heated on platinum foil; the sulphite and carbonate are decomposed with effervescence by acids | Borates. Oxalates.<br>Sulphites. Tartrates.<br>Sulphates. Phosphates.<br>Carbonates. Citrates.<br>Of these white calcium precipitates, the sulphate only is sol. in much water; the borate, carbonate, and citrate are sol. in solution of chloride of ammonium; all are sol. in acetic acid except oxalate and some tartrate and sulphate; all are sol. in hydrochloric acid, much sulphate excepted; the dry tartrate and citrate char when heated; the sulphite and carbonate effervesce with acids. | Chlorides, white.<br>Bromides, white.<br>Iodides, yellow.<br>Cyanides, white.<br>Borates, white.<br>Sulphides, black.<br>Sulphites, white.<br>Carbonates, white.<br>Oxalates, white.<br>Tartrates, white.<br>Phosphates, yellow.<br>Citrates, white.<br>Of these silver precipitates, the chloride, bromide, iodide, cyanide, and sulphide are insoluble in dilute nitric acid; the rest soluble. | Borates, yellowish.<br>Sulphides, black.<br>Carbonates, reddish.<br>Oxalates, yellow.<br>Phosphates, yel.-white.<br>Gives red color with acetates, if neutral. | Nitrates.<br>Chlorates.<br>Apply special tests. |

*Note.*—The student should practise the examination of aqueous solutions of salts until able to detect acidulous radicals with facility and precision. For this purpose he may finish the analyses of salts or solutions already examined for common and rarer metals, or have aqueous solutions of salts, or the salts themselves, specially prepared for present use, the metals of the salts being stated. He will then be in a position to effectively study the analysis of salts which may or may not be soluble in water, examining them for both basylous and acidulous radicals.

### REMARKS ON THE PRECEDING TABLE.

The first point of value to be noticed in connection with this Table is one of a negative character; namely, if either of the reagents gives no reaction, it is self-evident that the salts which it decomposes with production of a precipitate must be absent. Then, again, if the action of one of the reagents indicates the absence of certain acidulous radicals, those radicals cannot be precipitated by the other reagents; thus, if the action of sulphuric acid points to the absence of sulphides, sulphites, carbonates, cyanides and acetates, these salts may be struck out of the other lists, and the examination of subsequent precipitates be so far simplified. Or, if the barium precipitate is soluble in hydrochloric acid and the calcium precipitate in acetic acid, neither sulphates nor oxalates can be present. Observing these and other points of difference, which will be seen on careful and thoughtful reflection, and remembering the facts suggested by a knowledge of what basylous radicals are present, one acidulous radical after the other may be struck off as absent or present, leaving only one or two as the objects of special experiment. Among the chief difficulties to be encountered will be the separation from each other of chlorides, bromides, iodides, and cyanides, or of tartrates from citrates, and confirmatory tests of the presence of certain compounds. These may all be surmounted on referring back to the reactions of the various radicals, as described under their hydrogen salts, the acids.

In rendering a solution neutral, for the application of the various group-tests, the employment of any large amount of acid or of alkali must be noted; the presence of actual alkalies (that is *hydrates*) or of acids, respectively, being thereby indicated.

Sulphuric acid, the first group-test, may itself yield, especially when heated with some solid substances, sulphurous acid or hydrosulphuric acid (see pp. 306 and 311); hence the production of the latter acids from a *diluted* solution only is evidence of the presence of a sulphide or sulphite.

In the precipitate produced by chloride of barium, the second group-test, the oxalic radical may be specially sought by the test described in the "note" on p. 317.

Chloride of calcium does not precipitate citrates readily or completely in the cold, therefore the mixture should be filtered and the filtrate boiled; calcium citrate then falls. Calcium tartrate is soluble in solution of chloride of ammonium when quite freshly precipitated, but not after it has become crystalline. From their solution in chloride of ammonium, carbonate of calcium is mostly precipitated by ammonia, and citrate on boiling.

The rarer acidulous radicals will very seldom be met with. *Benzoates*, *hippurates* (which give benzoic acid), *hypochlorites*, *hyposulphites*, *nitrites*, and *valerianates* show themselves under the sulphuric treatment. *Ferrocyanides*, *ferridcyanides*, *meconates*, *succinates*, *sulphocyanates*, *tannates*, and *gallates* appear among the salts whose presence is indicated by ferric chloride; *formiates*, *hypophosphites*, *malates*, and others by nitrate of silver. *Urates* char when heated, giving an odor resembling that of burnt feathers.

In actual practice the analyst nearly always has some clue to the nature of rarer substances placed in his hands.

If chromium and arsenicum have been detected among the basylous radicals, those elements may be present in the form of *chromates*, *arseniates*, and *arsenites*, yielding with chloride of barium yellow chromate of barium and white arseniate and arsenite of barium, and with nitrate of silver red chromate, brown arseniate, and yellow arsenite of silver.

---

## QUESTIONS AND EXERCISES.

642. In analyzing an aqueous solution of salts, for which radicals would you first search, the basylous or the acidulous? and why?

643. In an aqueous solution there have been found magnesium (Mg) and potassium (K), with the sulphuric radical ($SO_4$) and iodine (I); state the nature of the salts which were originally dissolved in the water, and mention the principles which guide you in the conclusions.

644. Give a sketch of the method by which to analyze a neutral or only faintly acid aqueous liquid for the acidulous radical of salts. In what stage of the process would the following salts be detected?

*a*. Carbonates and Sulphates.
*b*. Oxalates.
*c*. Tartrates and Nitrates.
*d*. Acetates and Sulphites.
*e*. Bromides and Cyanides.
*f*. Borates.
*g*. Iodides and Phosphates.
*h*. Chlorates, Oxalates, and Acetates.
*i*. Chlorides and Iodides.
*j*. Sulphites.
*k*. Sulphides, Carbonates, and Nitrates.
*l*. Citrates and Sulphates.

645. Nitrate of silver gives no precipitate in an aqueous solution; what acidulous radicals may be present?

646. Chloride of barium gives no precipitate in a neutral solution, but nitrate of silver a white; what acidulous radicals are indicated?

647. Ferric chloride produces a deep red color in a solution, chloride of calcium yielding no precipitate; what salts may be present, and how may they be distinguished from each other?

648. Ferric chloride gives a black precipitate in a solution in which sulphuric acid develops no odor; to what is the effect due?

# ANALYSIS OF SALTS.

## SINGLE OR MIXED, SOLUBLE OR INSOLUBLE.

Thus far all material substances, especially those of pharmaceutical interest, have been regarded as being definite compounds, and as having certain well-defined parts, termed, for convenience, basylous and acidulous respectively; moreover attention has been designedly restricted to those definite compounds which are soluble in water. But there are many substances having no definite or known composition; and of those having definite composition there are many having no definite or ascertained parts. Again, of those having definite composition, and whose constitution admits of the entertainment of theory, there are many insoluble in water.

Chemical substances of whose composition or constitution little or nothing is at present known, are chiefly of animal and vegetable origin, and figure in tables of analysis under the convenient collective title of "extractive matter;" they are not of immediate importance, and may be omitted.

Of substances which are definite in composition, but whose parts or radicals, if they have any, are unknown or imperfectly known, there are only a few (such as the alkaloids, amylaceous and saccharine matters, the glucosides, alcoholic bodies, albuminoid, fatty, resinoid, and colorific substances) which have any considerable amount of medical or pharmaceutical interest; these will be noticed subsequently.

Definite compounds most frequently present themselves; and of these by far the larger proportion (namely, the salts soluble in water) have already been fully studied. There remain, however, many salts which are insoluble in water, but which must be brought into a state of solution before they can be effectively examined from any analytical, pharmaceutical, or a physiological point of view. The next subject of laboratory work is therefore the analysis of substances which may or may not be soluble in water. This will involve no other analytical schemes than those which have been given, will in only one or two cases increase the difficulty of the analysis of the precipitate produced by a group-reagent, but will give roundness, completeness, and a practical bearing to the reader's analytical knowledge. Such a procedure will at the same time bring into notice the methods by which substances insoluble in water are manipulated for pharmaceutical purposes, or made available for use as food by plants, or as food and medicine by man and animals generally.

### *Preliminary Examination of Solid (chiefly mineral) Salts.*

Before attempting to dissolve a salt for analysis, its appearance and other physical properties should be noted, and the influence of heat and strong sulphuric acid be ascertained. If the operator knows how to interpret what is thus observed, and to what extent to place

confidence in the observations, he may more certainly obtain a high degree of precision in analysis, and will always gain some valuable negative information. But if he has only slight experience of the appearance and general properties of bodies, or has the habit of turning what should be inferences from tentative processes into foregone conclusions, he should omit the preliminary examination altogether, or only follow it out under the guidance of a judicious tutor; for it is impracticable here to do more than hint at the results which may be obtained by such an examination, or to so adapt description as to prevent a student from allowing unnecessary weight to preconceived ideas.

Whatever be the course pursued, short memoranda describing results should invariably be entered in the note-book.

1. Examine the physical characters of the salt in various ways, but never, or only rarely, by the palate, on account of the danger to be apprehended.

If the salt is white, colored substances cannot be present; if colored, the tint may indicate the nature of the substance or of one of its constituents, supposing that the learner is already acquainted with the colors of salts. Closer observation, aided perhaps by a lens, may reveal the occurrence, in a pulverulent mixture, of small crystals or pieces of a single substance; these should be picked out by a needle and examined separately. In a powder or roughly divided mixture of substances, the process of sifting (through such sieves as muslin of different degrees of fineness) often mechanically separates substances, and thus greatly facilitates analysis. The body may present an undoubted metallic appearance, in which case only the metals existing under ordinary atmospheric conditions need be sought. Peculiarity in smell reveals the presence of ammonia, hydrocyanic acid, hydrosulphuric acid, etc. Between the fingers a substance is, perhaps, hard, soft, or gritty; consequent inferences follow. Or the matter may be heavy, like the salts of barium or lead; or light, like the carbonates and hydrates of magnesium; or may be one of the pharmaceutically well-known class of "scale" preparations.

2. Place a grain or two of the salt in a small dry test-tube or in a piece of ordinary tubing, closed at one end, and heat it, at first gently, then more strongly, and finally, if necessary, by the blowpipe.

Gases or vapors of characteristic appearance or odor may be evolved; such as iodine, nitrous fumes, sulphurous, hydrocyanic, or ammoniacal gases. Much steam given by a dry substance indicates either hydrates or salts containing water of crystallization. (A small quantity of interstitial moisture often causes heated crystalline substances to *decrepitate*—from *decrepo*, I crackle—that is, break up with slight explosive violence, owing to the expansive force of the steam suddenly generated.) A sublimate may be obtained, due to salts of mercury or arsenicum, to oxalic or benzoic acid, or to sul-

phur free or as a sulphide—a salt wholly volatile containing such substances only. The compound may blacken, pointing to the presence of organic matter—which, in common definite salts, will probably be in the form of acetates, tartrates, and citrates, or as common salts of the alkaloids morphia, quinia, strychnia, or as starch, sugar, salicin, or in other definite or indefinite forms common in pharmacy and for which tests will be given in subsequent pages. If no charring occurs, the important fact that no organic matter is present is established. The residue may change color from presence or development of oxide of zinc, oxide of iron, etc., or melt from the presence of a fusible salt and absence of any large proportion of infusible salts, or being unaltered, showing the absence of any large amount of such substances.

3. Place a grain or two of the salt in a test-tube, add a drop or two of strong sulphuric acid, cautiously smelling any gas that may be evolved; afterward slowly heat the mixture, noticing the effect, and stopping the experiment when any sulphuric fumes begin to escape.

Iodine, bromine, and nitrous or chlorinoid fumes will reveal themselves by their color, indicating the presence of iodides, bromides, iodates, bromates, nitrates, and chlorates. The evolution of a colorless gas fuming on coming into contact with air, and having an irritating odor, points to chlorides, fluorides, or nitrates. Gaseous products having a greenish color and odor of chlorine indicate chlorates, hypochlorites, or chlorides mixed with other substances. Slight sharp explosions betoken chlorates. Evolution of colorless gas may proceed from cyanides, acetates, sulphides, sulphites, carbonates, or oxalates. Charring will be due to citrates, tartrates, or other organic matter. If none of these effects are produced, most of the bodies are absent or only present in minute quantity. The substances apparently unaffected by the treatment are metallic oxides, borates, sulphates, and phosphates.

4. Exposure of the substance to the blowpipe-flame, on platinum wire with or without a bead of borax or microcosmic salt (phosphate of sodium, ammonium, and hydrogen, $NaAmHPO_4$) —on platinum foil, in a porcelain crucible, or on a crucible lid with or without carbonate of sodium—on charcoal, alone or in conjunction with carbonate of sodium, cyanide of potassium, or nitrate of cobalt, will sometimes yield important information, especially to one who has devoted much attention to reactions producible by the blowpipe-flame. The medical or pharmaceutical student, however, will seldom have time to work out this subject to an extent sufficient to make it a trustworthy guide in analysis. (See Plattner and Muspratt "On the Use of the Blowpipe," and a chapter in Galloway's "Manual of Quantitative Analysis.")

*Methods of dissolving and analyzing single or mixed solid substances.*

*Having submitted the substance to preliminary examination, proceed to dissolve and analyze by the following methods. These operations consist in treating a substance well powdered, consecutively with cold or hot water, hydrochloric acid, nitric acid, nitro-hydrochloric acid, or fusion with alkaline carbonates and solution of the product in water and acid. Resulting liquids are analyzed in the manner already described, or by slightly modified processes as detailed in the following paragraphs.*

*Solution in Water.*—Boil about a grain of the salt presented for analysis in about a third of a test-tubeful of water. If it dissolves prepare a solution of about 20 or 30 grains in half an ounce or more of water, and *proceed with the analysis in the usual way*, testing first for the basylous radical or radicals by the proper group reagents (HCl, $H_2S$, AmHS, $Am_2CO_3$, $Am_2HPO_4$), p. 224 or 259, and then for the acidulous radical or radicals, directly or by aid of the prescribed reagents ($H_2SO_4$, $BaCl_2$, $CaCl_2$, $AgNO_3$, $Fe_2Cl_6$), p. 366.

If the salt is not *wholly* dissolved by the water, ascertain whether or not any has entered into solution, by filtering, if necessary, and evaporating a drop or two of the clear liquid to dryness on platinum foil; the presence or absence of a residue gives the information sought. If anything is dissolved, prepare a sufficient quantity of solution for analysis and proceed as usual, reserving the insoluble portion of the mixture, after thoroughly exhausting with water, for subsequent treatment by acids.

*Solution in Hydrochloric Acid.*—If the salt is insoluble in water, digest about a grain of it (or of the insoluble portion of a mixed salt) in a few drops of hydrochloric acid, adding water, and boiling if necessary. If the salt wholly dissolves, prepare a sufficient quantity of the liquid, noticing whether or not any effervescence (due to the presence of sulphides, sulphites, carbonates, or cyanides) occurs, and proceed with the analysis as before, except that the first step, the addition of hydrochloric acid, may be omitted.

The analysis of this solution will in most respects be simpler than that of an aqueous solution, inasmuch as the majority of salts (all those soluble in water) will be absent. This acid solution will, in short, only contain: chlorides produced by the action of the hydrochloric acid on sulphides, sulphites, carbonates, cyanides, oxides, and hydrates; and certain borates, oxalates, phosphates, tartrates, and citrates (possibly silicates and fluorides) which are insoluble in water but soluble in acids without apparent decomposition. The first four

—sulphides, sulphites, carbonates, and cyanides—will have revealed themselves by the occurrence of effervescence during solution; and the presence of oxides and hydrates may often be inferred by the absence of compatible acidulous radicals. The borates, oxalates, phosphates, tartrates, and citrates alluded to will be reprecipitated in the general analysis as soon as the acid of the solution is neutralized: that is, will come down in their original state when ammonia and sulphydrate of ammonium are added to the usual course. Of these precipitates, only the oxalate of calcium and the phosphates of calcium and magnesium need occupy attention now; for oxalate and phosphate of barium seldom or never occur, and the borates, tartrates, and citrates met with in medicine or in general analysis are all soluble in water. These phosphates and oxalates, then, will be precipitated in the course of analysis along with iron, their presence not interfering with the detection of any other metal. If, from the unusual light color of the ferric precipitate, phosphates and oxalates are suspected, the precipitate is treated according to the following Table (reference to which should be inserted in the Table for metals, under Fe, pp. 223 and 258).

PRECIPITATE OF PHOSPHATES, OXALATES, AND FERRIC HYDRATE.

Dissolve in HCl, add citric acid, then $NH_4HO$, and filter.

<table>
<tr><td rowspan="3">Filtrate<br>Fe.<br>Add HCl and $K_4Fcy$.<br>Blue ppt.</td><td colspan="3">Precipitate<br>$Ca_3 2PO_4$, $CaC_2O_4$, $Mg_3 2PO_4$.<br>Boil in acetic acid and filter.</td></tr>
<tr><td>Insoluble<br>$CaC_2O_4$.<br>White.</td><td colspan="2">Filtrate<br>$Ca_3 2PO_4$, $Mg_3 2PO_4$.<br>Add $Am_2C_2O_4$, stir, filter.</td></tr>
<tr><td></td><td>Precipitate<br>white, including<br>$Ca_3 2PO_3$.</td><td>Filtrate.<br>Add AmHO.<br>White ppt.<br>$MgNH_4PO_4$.</td></tr>
</table>

In analyzing phosphates and oxalates, advantage is also frequently taken of the facts that the phosphoric radical is wholly removed from solution of phosphates in acid by the addition of an alkaline acetate, ferric chloride, and subsequent ebullition, as described under "Phosphoric Acid" (p. 331), and that dry oxalates are converted into carbonates by heat, as mentioned under "Oxalic Acid," (p. 317). See also p. 331, 4th Analytical Reaction.

Certain arseniates and arsenites, insoluble in water but soluble in hydrochloric acid, may accompany the above phosphates and oxalates

if from any cause hydrosulphuric acid gas has not been previously passed through the solution, or passed for an insufficient length of time.

If the substance insoluble in water does not *wholly* dissolve in hydrochloric acid, ascertain if any has entered into solution, by filtering, if necessary, and evaporating a drop of the clear liquid to dryness on platinum foil; the presence or absence of a residue gives the information sought. If anything is dissolved, prepare a sufficient quantity of solution for analysis, and proceed as usual, reserving the insoluble portion of the mixture, after thoroughly exhausting with hydrochloric acid and well washing with water, for the following treatment by nitric acid.

*Solution in Nitric Acid.*—If the salt is insoluble in water and hydrochloric acid, boil it (or that part of it which is insoluble in those menstrua) in a few drops of nitric acid. If it wholly dissolves, remove excess of acid by evaporation, dilute with water, and proceed with the analysis.

This nitric solution can contain only very few substances; for nearly all salts soluble in nitric acid are also soluble in hydrochloric acid, and therefere will have been previously removed. Some of the metals, however (Ag, Cu, Hg, Pb, Bi), as well as amalgams and alloys, unaffected or scarcely affected by hydrochloric acid, are readily attacked and dissolved by nitric acid. Many of the sulphides, also insoluble in hydrochloric acid, are dissolved by nitric acid, usually with separation of sulphur. Calomel is converted, by long boiling with nitric acid, into mercuric chloride and nitrate. The nitrates here produced are soluble in water.

This nitric solution, as well as the hydrochloric and aqueous solutions, should be examined separately. Apparently time would be saved by mixing the three solutions together and making one analysis. But the object of the analyst is to separate every radical from every other; and when this has been partially accomplished by solvents, it would be unwise to again mix and separate a second time. Moreover, solvents often do what the chemical reagents cannot—namely, separate *salts* from each other. This is important, inasmuch as the end to be obtained in an analysis is not only an enumeration of the radicals present, but a statement of the actual condition in which they are present; the analyst must, if possible, state of what salts a given mixture was originally formed—how the basylous and acidulous radicals were originally distributed. In attempting this, much must be left to theoretical considerations; but a process by which the salts themselves are separated is of trustworthy practical assistance; hence the chief advantage of analyzing separately the solutions resulting from the action of water and acids on a solid substance.

*Solution in Nitro-Hydrochloric Acid.*—If the salt or any part of a mixture of salts is insoluble in water, hydrochloric acid, and nitric acid, digest it in nitro-hydrochloric acid,

warming or even boiling gently, if necessary; evaporate to remove excess of acid, dilute, and proceed as before.

Sulphide of mercury and substances only slowly attacked by hydrochloric or nitric acid, as for example, calomel and ignited ferric oxide, are sufficiently altered by the free chlorine of aqua regia to become soluble.

*If the substance is insoluble in water and acids*, it is one or more of the following substances: Sand and certain silicates, such as pipeclay and other clays; fluor spar; cryolite ($3NaF, AlF_3$); sulphates of barium, strontium, and possibly calcium; tinstone; glass; felspar (double silicate of aluminium and other metals); chloride of silver; sulphate of lead. It may also be or contain carbon or carbonaceous matter, in which case it is black and combustible, burning entirely or partially away when heated in the air—or be or contain sulphur, in which case sulphurous gas is evolved, detected by its odor, when the substance is exposed to heat. For the other substances proceed according to the following (Bloxam's) method:—

Four or five grains of the substance are intimately mixed with twice the quantity of dried carbonate of sodium, and this mixture well rubbed in a mortar with five times its weight of *deflagrating flux* (1 of finely powdered charcoal to 6 of nitre). The resulting powder is placed in a thin porcelain dish, or crucible, or clean iron tray, and a lighted match applied to the centre of the heap. Deflagration ensues, and decomposition of the various substances occurs, the acidulous radicals going to the alkali-metals to form salts soluble in water, the basylous radicals being simultaneously converted into carbonates or oxides. The mass is boiled in water for a few minutes, the mixture filtered, and the residue well washed. The filtrate may then be examined for acidulous radicals and aluminium, and the residue dissolved in dilute hydrochloric acid and analyzed by the ordinary method.

The only substance which resists this treatment is chrome-iron ore.

To detect alkali in felspar, glass, or cryolite, Bloxam recommends deflagration of the powdered mineral with one part of sulphur and six of nitrate of barium. The mass is boiled in water, the mixture filtered, hydrate and carbonate of ammonium added to remove barium, the mixture again filtered, and the filtrate evaporated and examined for alkalies by the usual process.

### *Hydrates and Oxides.*

If no acidulous radical can be detected in a substance under analytical examination, or if the amount found is obviously in-

sufficient to saturate the quantity of basylous radical present, the occurrence of oxides or hydrates, or both, may be suspected. Confirmation of their presence will be found in the general rather than in any special behavior of the substances. Some hydrates yield water when heated—in a dry test-tube held nearly horizontally in a flame, so that moisture may condense on the cool part of the tube. Some oxides yield oxygen—detected by heating in a test-tube, and inserting the incandescent end of a strip of wood. Soluble hydrates cause abundant evolution of ammonia gas when heated with solution of chloride of ammonium. Soluble hydrates also give characteristic precipitates with the various metallic solutions. Hydrates and oxides insoluble in water not only neutralize much nitric acid or acetic acid, but are thereby converted into salts soluble in water. Most oxides and hydrates have a characteristic appearance. In short, some one or more properties of an oxide or a hydrate will generally betray its presence to the student who not only has knowledge respecting chemical substances, but has cultivated the faculties of observation and perception.

---

## QUESTIONS AND EXERCISES.

649. Describe the preliminary treatment to which a salt may be subjected prior to systematic analysis.

650. Mention substances which might be recognized by smell.

651. Which classes of salts are heavy, and which light?

652. Name some bodies detectable by their color.

653. What inference may be drawn from the appearance of steam when dry substances are heated?

654. Why do certain crystals decrepitate?

655. If a powder sublimes on being heated, to what classes of compounds may it belong?

656. When heat causes charring, what conclusion is drawn?

657. No change occurring by heat, which substances cannot be present?

658. Give example of salts which are identified by their reaction with strong sulphuric acid; and by their comportment in the blowpipe flame, with or without borax or microcosmic salt.

659. What are the solvents usually employed in endeavoring to obtain a substance in a state of solution, and what is the order of their application?

660. Name a few salts which may be present in an aqueous solution.

661. Mention some common compounds insoluble in water, but soluble in hydrochloric acid.

662. What substances are attacked only by nitric acid or nitrohydrochloric acid?

663. At what stage of analysis do arsenites and arseniates show themselves?

664. Sketch out a method for the complete analysis of a liquid suspected to be an aqueous solution of neutral salts.

665. How can earthy phosphates and oxalates with ferric oxide be separated from each other?

666. How would you proceed to analyze an alloy?

667. By what process may substances insoluble in water or acids be analyzed?

668. How would you qualitatively analyze glass?

---

## RECAPITULATORY AND OTHER NOTES ON THE CONSTITUTION OF THE DEFINITE CHEMICAL COMPOUNDS COMMONLY TERMED SALTS.

The molecules of a salt contain radicals—which may be either elementary or compound: pp. 38 and 66.

Each radical has a definite exchangeable value: p. 121.

The definite exchangeable values of radicals differ in different series of radicals: pp. 122, 123.

In one and the same molecule of a salt, two or more different atoms of the same element may possess the two distinct functions of being (*a*) a single definite distinct radical and (*b*) one member of a group of atoms which together form a single definite distinct radical: p. 264.

The relation to each other, either of the elementary or the compound radicals, in organic substances or salts, is apparently far more complex than the relation to each other of the elementary and compound radicals in inorganic salts: pp. 379 and 380.

The properties of salts are regarded as depending on (*a*) the nature, (*b*) the number, and (*c*) the position in relation to each other, of the elementary and compound radicals in a molecule.

Dumas, afterwards Laurent, and then Gerhardt, attempted the classification of salts under such *types* as the following:—

| The hydrogen type. | The water type. | The ammonia type. |
|---|---|---|
| $\left.\begin{matrix}H\\H\end{matrix}\right\}$ | $\left.\begin{matrix}H\\H\end{matrix}\right\}O$ | $\left.\begin{matrix}H\\H\\H\end{matrix}\right\}N$ |

Other chemists have extended the number of such types of salts. Further, by writing the typical formulæ in the above and other manners, a mode of indicating the facts assumed to be dependent on the position of the atoms in a molecule has been sought to be obtained. Finally, the natural development of this train of thought and of practice has produced the *graphic formulæ* of Kékulé, Frankland, and others.

*Caution.*—The conjectural or theoretic character of our ideas respecting masses of matter being formed of molecules, and molecules of atoms, and that molecules contain radicals consisting of one or more atoms, must never be lost sight of—highly valuable and prac-

tically useful though the hypotheses be: pp. 38, 51, 264, 265, 286, 300.

*Berthollet's Laws.*—"When we cause two salts to react by means of a solvent, if, in the course of double decomposition, a new salt can be produced less soluble than those which we have mixed, this salt will be produced. When we apply dry heat to two salts, if, by double decomposition, a new salt can be produced more volatile than the salts previously mixed, this salt will be produced."

*Malaguti's Law.*—When solutions of two different salts are mixed and metathesis occurs and four salts result, the proportions of the salts to each other are dependent on the strength or intensity of force with which the respective basylous and acidulous radicals are united.

The state of equilibrium just mentioned may be permanent or temporary. The latter condition obtains when one of the salts which may possibly be produced is insoluble, for as soon as precipitation occurs the equilibrium is upset, and is re-established only to be upset again, and so on until from the four salts there result one in solution and one out of solution. This would seem to be the way in which the laws termed Berthollet's work.

# CHEMISTRY OF CERTAIN SUBSTANCES OF VEGETABLE AND ANIMAL ORIGIN.

Except alcohol and a few acids, the compounds which have hitherto engaged notice have been of mineral origin. But the two other kingdoms of nature, the animal and vegetable, furnish a large number of definite substances. These, indeed, when discovered, were producible only by organized living structures, and were hence termed *organic* compounds.*

A few of these compounds of common occurrence in pharmacy and possessing prominent characteristics, may now occupy attention; reactions of the alkaloids and some other principles may be performed, and the methods of examining morbid urine be experimentally studied. There will then remain to be studied certain galenical, as distinguished from chemical, substances, solid and liquid, which can only be fairly regarded from a pharmacist's rather than a chemist's point of view, and a still larger number, doubtless, not yet brought within the grasp of chemist or pharmacist, and of which, therefore,

* *Organic*, from ἄ,γανω, *organon*, an organ. A large number of organic compounds can now be obtained artificially—without the aid of a living organism; hence the distinction formerly drawn between organic and inorganic compounds, organic and inorganic chemistry, is fast breaking down.

we must at present be content to remain in ignorance. An opportunity, however, will be afforded of noticing the effect of such organic matter as a vomit or the contents of a stomach, in masking or preventing the reactions by which mineral and vegetable poisons are detected.

---

## ALKALOIDS.

### *Constitution of Alkaloids or Organic Bases.*

*Natural Alkaloids.*—The alkaloids, or alkali-like bodies (*εἶδος, eidos*, likeness) have many analogies with ammonia. Their constitution is not yet known; but they are probably derivatives of a single molecule of ammonia ($NH_3$), or of double, triple, or quadruple molecules ($N_2H_6$, $N_3H_9$, $N_4H_{12}$).

*Artificial Alkaloids.*—Numerous artificial alkaloids or organic bases, unquestionably having the constitution just mentioned, have already been formed. These are sometimes termed *amines*, and are primary, secondary, and tertiary, according as one, two, or three atoms of hydrogen in ammonia have been displaced by radicals, as seen in the following general formulæ (R = any univalent radical. *Vide* Index, "Alcohol radicals")—

$$\left.\begin{matrix} R \\ H \\ H \end{matrix}\right\} N \qquad \left.\begin{matrix} R \\ R \\ H \end{matrix}\right\} N \qquad \left.\begin{matrix} R \\ R \\ R \end{matrix}\right\} N$$

or the following examples—

$$\left.\begin{matrix} C_2H_5 \\ H \\ H \end{matrix}\right\} N \qquad \left.\begin{matrix} C_2H_5 \\ C_2H_5 \\ H \end{matrix}\right\} N \qquad \left.\begin{matrix} C_2H_5 \\ C_2H_5 \\ C_2H_5 \end{matrix}\right\} N$$

Ethylamine or ethylia ($C_2H_7N$). Diethylamine or diethylia ($C_4H_{11}N$). Triethylamine or tryethylia ($C_6H_{15}N$).

The three classes have also been termed amidogen-, imidogen-, and nitrile-bases.

*Mode of formation of Artificial Alkaloids.*—A few illustrations will suffice. Just as the addition of iodide of hydrogen (HI) to ammonia (that is, the common tri-hydrogen ammonia, $NH_3$) gives iodide of common ammonium (NHHHHI or $NH_4I$), so the addition of iodide of ethyl ($C_2H_5I$ or EtI) (see Index) to ammonia ($NH_3$) gives the iodide of ethyl-ammonium (NHHHEtI, or $NH_3EtI$, or $NH_3C_2H_5I$). A fixed alkali turns out common *ammonia* (NHHH) from the iodide (or any other salt) of common ammonium; it turns out ethyl-ammonia (NHHEt) from the iodide (or any other salt) of ethyl-ammonium. Ethyl-ammonia (or *ethylia*, or ethylamine), NHHEt, with iodide of ethyl, EtI, gives iodide of diethyl-ammonium (NHHEtEtI, or $NH_2Et_2I$, or $NH_2[C_2H_5]_2I$). From the latter, potash turns out diethyl-ammonia ($NHEt_2$); diethyl-ammonia (*diethylia* or diethylamine) with iodide of ethyl gives iodide of triethyl-ammonium ($NHEt_3I$). The latter with alkali gives triethyl-ammonia or *triethylia* or triethylamine ($NEt_3$), and this with iodide of ethyl gives iodide of tetrethyl-ammonium, $NEt_4I$.

What has just been stated respecting iodide of ethyl is true of other salts of ethyl; and what is true of salts of ethyl is true of salts of an immense number of other radicals—univalent, bivalent, etc.: so that a vast number of artificial alkaloids and their salts can be produced. The reactions are not always so sharp as those just given. Mixtures of primary, secondary, and tertiary compounds rather than either alone often result in an experiment; but the reactions are typically true.

Some of these artificial alkaloids not only resemble natural alkaloids but are strong caustic liquids, like solution of ammonia.

Then, the displacing radical in an artificial alkaloid or its salt may not only be of one kind, as indicated in the preceding paragraphs, but of different kinds; and while the radical displacing one atom of hydrogen is keeping its place, any of the many known radicals may occupy the position of one or all of the other atoms of hydrogen. Thus, for example, we have methyl-ethyl-amyl-amine ($C_8H_{19}N$, or $CH_3C_2H_5C_5H_{11}N$, or MeEtAyN), a colorless oily body, of agreeable aromatic odor. The empirical formulæ of morphia, quinia, etc., may some day be similarly resolvable into rational formulæ. Their artificial production will then quickly follow.

*Propylamine* or *tritylia* ($C_3H_7HHN$) is a volatile oil, one product of the destructive distillation of bones and other animal matters.

The organic bases derived from one molecule of ammonia are termed monamines; from two molecules, diamines; from three, triamines; and from four, tetramines:—

$$\left.\begin{matrix}R\\R\\R\end{matrix}\right\}N \qquad \left.\begin{matrix}R_2\\R_2\\R_2\end{matrix}\right\}N_2 \qquad \left.\begin{matrix}R_3\\R_3\\R_3\end{matrix}\right\}N_3 \qquad \left.\begin{matrix}R_4\\R_4\\R_4\end{matrix}\right\}N_4$$

In these amines any bivalent, trivalent, or quadrivalent radical may occupy the place of two, three, or four univalent radicals.

*Evidence of Constitution of the Natural Alkaloids.*—Attempts to form artificially the important natural organic bases have hitherto failed; but the primary, secondary, or tertiary character of some of them has been indicated by the introduction or elimination of methyl, ethyl, and other radicals for hydrogen.

*Note on Nomenclature of Natural Alkaloids.*—The first syllables of the names of the natural alkaloids recall the name of the plant whence they were obtained, or some characteristic property. It is to be regretted that the last syllable is not either *ine* or *ia*, instead of sometimes one and sometimes the other: general usage seems to be in favor of the latter, a plan that distinguishes the alkaloids from some other substances the names of which end in *ine*, as chlorine, bromine, iodine, fluorine, glycerine, gelatine, etc. The names of the salts of the alkaloids are given on the assumption that the acid unites with the alkaloid without decomposition. Thus hydrochlorate of morphia is regarded as morphia with hydrochloric acid, just as we might assume sal-ammoniac to be ammonia ($NH_3$) with hydrochloric acid (HCl), and name it hydrochlorate of ammonia ($NH_3HCl$) instead of chloride of ammonium ($NH_4Cl$). All acids, even sulphydric, unite with alkaloids and form salts having similar names.

*Antidotes.*—In cases of poisoning by alkaloids, emetics and the

stomach-pump must be relied on rather than chemical agents. But astringent liquids may be administered, for tannic acid precipitates many of the alkaloids from their aqueous solution, absorption of the poison being thus possibly retarded.

---

## MORPHIA, OR MORPHINE.

Formula, $C_{17}H_{19}NO_3.H_2O$. Molecular weight 303.

*Occurrence.*—Morphia occurs in opium (the inspissated juice of the fruit, *Papaveris capsulæ*, of the White Poppy, *Papaver somniferum*) as meconate of morphia The dried poppy-capsule of pharmacy contains opium principles, but varying much in nature and proportion. Ordinary Asia Minor opium (Turkey, Smyrna, or Constantinople opium) contains, when dried, from 10 to 15 per cent. of morphia.

*Morphia*, U. S. P., is made by adding to infusion of opium an equal bulk of alcohol, then slight excess of ammonia, and setting aside for crystalline morphia to separate. It is purified by recrystallization.

*Process for Hydrochlorate.*—The hydrochlorate, $C_{17}H_{19}NO_3,HCl,3H_2O$ (*Morphiæ Hydrochloras*, B. P.), occurs in slender white acicular crystals; it is prepared by simply decomposing an aqueous infusion of opium with chloride of calcium, meconate of calcium and hydrochlorate of morphia being produced. (If the infusion, which is always acid, be first nearly neutralized by the cautious addition of small quantities of very dilute solution of ammonia, the chloride of calcium then at once causes a precipitate of meconate of calcium, which can be filtered off, leaving a colored solution of hydrochlorate of morphia. On the large scale—*vide* B. P.—the details are somewhat different.) The salt is partially purified by crystallization from the evaporated liquid, then by treatment of the solution of the impure hydrochlorate by animal charcoal, and lastly by precipitation of the morphia from the still colored liquid by ammonia and resolution of the morphia in hot dilute hydrochloric acid; hydrochlorate of morphia separates out on cooling. The process of U. S. P. consists in neutralizing morphia by hydrochloric acid. (*Morphiæ Murias*, U. S. P.)

*Morphiæ Sulphas*, U. S. P., by neutralizing morphia with sulphuric acid.

*Process for Acetate.*—Acetate of morphia ($C_{17}H_{19}NO_3C_2H_4O_2$) (*Morphiæ Acetas*, B. P. and U. S. P.) is a white pulverulent salt prepared by dissolving pure morphia in acetic acid, the morphia being prepared (B. P.) from a solution of the hydrochlorate by precipitation with ammonia. Eight parts of hydrochlorate yield about seven of acetate. One grain of acetate, so made, in twelve minims of water forms the *Injectio Morphiæ Hypodermica*, B. P.

Both the hydrochlorate and acetate of morphia are soluble in water, but the solution is not stable unless acidulated and containing alcohol; hence the official solutions, 4 grains in one ounce—1 in 110

(*Liquor Morphiæ Hydrochloratis*, B. P., and *Liquor Morphiæ Acetatis*, B. P.), consist of three parts water and one part rectified spirit, a few minims per ounce of hydrochloric or acetic acid being added. Even solid acetate of morphia is unstable, slowly dissociating into acetic acid and morphia, hence the acid odor of acetate of morphia. *Liquor Morphiæ Sulphatis*, U. S. P., is a solution in water, 1 in 456. The other official preparations in the British Pharmacopœia are *Suppositoria Morphiæ*, *Trochisci Morphiæ*, and *Trochisci Morphiæ et Ipecacuanhæ*.

When a simple but strong solution of acetate of morphia in water is required (for hypodermic injection) one part of the salt, if recently made, may be dissolved in six parts of water (1 grain in 11 grains, B. P.). If the acetate of morphia is old, it either will not dissolve, or, if dissolved by aid of warmth and a little acetic acid, will soon be re-deposited. The aqueous solution, however well made, cannot be kept in its normal condition for any great length of time.

*Other alkaloids* exist in opium. In the above process a considerable quantity of an alkaloid of very weak basic properties, *narcotine* ($C_{23}H_{23}NO_7$) (*Narcotina*, P. I.) remains in the exhausted opium, and may be extracted by digesting in acetic acid, filtering, precipitating by ammonia. It crystallizes in brilliant needles from alcohol or ether. By oxidation it yields *cotarnine* and an acid termed *opianic*. *Codeia* ($C_{18}H_{21}NO_3, H_2O$) is soluble in the slight excess of ammonia employed in precipitating the morphia. From the mother-liquors there have also been obtained *thebaia* ($C_{19}H_{21}NO_2$), *papaverine* ($C_{21}H_{21}NO_4$), *opianine* ($C_{21}H_{21}NO_7$?), *narceia* ($C_{22}H_{23}NO_9$), *cryptopia* ($C_{21}H_{23}NO_5$), *meconine* ($C_{10}H_{10}O_4$), *meconoisine* ($C_6H_{16}O_2$), *laudanine* ($C_{20}H_{23}NO_4$), *codamine* ($C_{20}H_{25}NO_4$), *gnoscopine* ($C_{34}H_{36}N_2O_{11}$), *pseudomorphia* ($C_{17}H_{19}NO_4$), *protopine* ($C_{20}H_{19}NO_5$), *laudanosine* ($C_{21}H_{27}NO_4$), *hydrocotarnine* ($C_{12}H_{15}NO_3$), *rhœadine* ($C_{20}H_{21}NO_6$), *meconidine* ($C_{21}H_{23}NO_4$), *lanthopine* ($C_{23}H_{25}NO_4$). A little acetic acid also exists in all opium (D. Brown).

### *Analytical Reactions.*

*First Analytical Reaction.*—To a minute fragment of a salt of morphia add one drop of water, and warm the mixture until the salt dissolves, then stir the liquid with a glass rod moistened by a strong *neutral* solution of perchloride of iron; a dirty-blue color is produced. This effect is not observed in dilute solutions.

*Second Analytical Reaction.*—To a drop or two of a strong solution of a morphia salt in a test-tube add a minute fragment of iodic acid ($HIO_3$, page 296); iodide is set free. Into the upper part of the tube insert a glass rod covered with mucilage of starch, and warm the solution; dark-blue starch iodide is produced. If the mixture of morphia and iodic acid be shaken up with chloroform or bisulphide of carbon, a violet solution is obtained.

This reaction is only confirmatory of others, as albuminous matters also reduce iodic acid.

*Third Analytical Reaction.*—To a few drops of an aqueous infusion of opium add a drop of *neutral* solution of perchloride of iron; a red solution of meconate of iron is produced. Add solution of corrosive sublimate; the color is not destroyed (as it is in the case of sulphocyanate of iron, a salt of similar tint).

In cases of poisoning by a preparation of opium, this test is almost as conclusive as a direct reaction of morphia (the poison itself), meconic acid being obtainable from opium only.

*Other Reactions.*—Add carbonate of sodium to a solution of a salt of morphia; a white precipitate of morphia falls, slowly and of a crystalline character if the solution is dilute. Collect this precipitate and moisten it with neutral solution of perchloride of iron; the bluish tint above referred to is produced. ——Add an alkali to a solution of hydrochlorate or acetate of the alkaloid; morphia is precipitated, soluble in excess of the fixed alkali, far less readily so in ammonia.——Moisten a particle of a morphia salt with nitric acid; an orange-red coloration is produced.——Heat morphia on platinum foil; it burns entirely away.

### *Apomorphia* ($C_{17}H_{17}NO_2$).

Apomorphia (ἀπό, *apo*, from, and *morphia*) is an alkaloid obtained from morphia by Matthiessen and Wright. It possesses remarkable physiological effects; one-tenth of a grain (in aqueous solution) injected under the skin, or one-quarter of a grain taken into the stomach, is said to produce vomiting in from four to ten minutes.

*Process.*—Hydrochlorate of morphia is hermetically sealed in a thick tube with considerable excess of hydrochloric acid, and heated to nearly 300° F. for two or three hours. The product is purified by diluting the contents of the tube with water, precipitating with bicarbonate of sodium, and treating the precipitate with ether or chloroform.——On shaking up the ethereal or chloroform solution with a very small quantity of strong hydrochloric acid, the sides of the vessel become covered with crystals of the hydrochlorate of the new base. These may be drained from the mother-liquor, washed with a little cold water, in which the salt is sparingly soluble, recrystallized from hot water, and dried on bibulous paper or over sulphuric acid. The formula ($C_{17}H_{17}NO_2$,HCl) indicates that the new alkaloid is derived from morphia by abstraction of the elements of water.

*Codeia*, also, according to the same chemists, yields apomorphia by similar treatment, a reaction that would seem to indicate that codeia is methyl-morphia:—

$$\underset{\text{Codeia.}}{C_{17}H_{17}CH_3HNO_3} + HCl = \underset{\text{Chl. of methyl.}}{CH_3Cl} + H_2O + \underset{\text{Apomorphia.}}{C_{17}H_{17}NO_2}$$

Dr. C. R. A. Wright has recently obtained several new derivatives of codeia.

## QUESTIONS AND EXERCISES.

669. Write some general formulæ of artificial alkaloids.

670. Name the substances represented by the following formulæ:—

$$\left.\begin{matrix} C_3H_7 \\ H \\ H \end{matrix}\right\} N, \quad \left.\begin{matrix} C_3H_7 \\ C_3H_7 \\ H \end{matrix}\right\} N, \quad \left.\begin{matrix} C\,H_3 \\ C_2H_5 \\ C_5H_{11} \end{matrix}\right\} N.$$

$$\left.\begin{matrix} CH_3 \\ H \\ H \end{matrix}\right\} N, \quad \left.\begin{matrix} CH_3 \\ CH_3 \\ H \end{matrix}\right\} N, \quad \left.\begin{matrix} CH_3 \\ CH_3 \\ CH_3 \end{matrix}\right\} N.$$

671. What is the assumed constitution of the salts of the alkaloids?

672. Describe the treatment in cases of poisoning by alkaloids.

673. Give the process for the preparation of Hydrochlorate of Morphia. In what form does morphia occur in opium?

674. How is Acetate of Morphia prepared?

675. What plan is adopted for preventing the decomposition of the official solutions of morphia?

676. Mention the analytical reactions of morphia.

677. In addition to the reactions of morphia, what test may be employed in searching for opium in a liquid or semifluid material?

678. How is apomorphia prepared, and what are its properties?

679. Describe the relation of morphia to codeia.

## QUINIA, OR QUININE.

Formula $C_{20}H_{24}N_2O_2,3H_2O$. Molecular weight 378

*Source.*—Quinia and other similar alkaloids exist in cinchona bark as kinates. In the yellow bark (*Cinchonæ Flavæ Cortex*, B. P. and U. S. P., from *Cinchona Calisaya*), chiefly quinia is present; in the pale bark (*Cinchonæ Pallidæ Cortex*, B. P. and U. S. P., chiefly from *C. Officinalis*), other alkaloids are more frequently found; while in the red bark (*Cinchonæ Rubræ Cortex*, B. P. and U. S. P.) these alkaloids occur in irregular proportions.

*Process for Sulphate.*—Sulphate of quinia (*Quiniæ Sulphas*, B. P.) is prepared according to the British Pharmacopœia by treating the yellow bark with dilute hydrochloric acid, precipitating the resulting solution of hydrochlorate of quinia by soda, and redissolving the precipitated quinia in the proper proportion of hot dilute sulphuric acid. This, the common commercial sulphate, crystallizes out on cooling in

silky acicular crystals, one molecule containing two atoms of quinia ($2C_{20}H_{24}N_2O_2$), one of sulphuric acid ($H_2SO_4$), and seven of water of crystallization ($7H_2O$).

In the process of the United States Pharmacopœia lime is used instead of soda, and the precipitated quinia is dissolved in boiling alcohol, the latter recovered by distillation, the residual quinia neutralized by diluted sulphuric acid, the solution treated with animal charcoal, filtered while hot, set aside to crystallize, and recrystallized if necessary.

Sulphate of quinia, the *common* or so-called *disulphate* ($(C_{20}H_{24}N_2O_2)_2,H_2SO_4,8H_2O$, is only slightly soluble in water; on the addition of dilute sulphuric acid, the so-called *neutral sulphate*, or *soluble sulphate* ($C_{20}H_{24}N_2O_4,H_2SO_4,7H_2O$), is formed, which is freely soluble. The latter salt may be obtained in large rectangular prisms.* An *acid sulphate* ($C_{20}H_{24}N_2O_2,2H_2SO_4,7H_2O$) also exists.

The ordinary disulphate of quinia is more soluble in alcohol or alcoholic liquids than in water; hence the *Tinctura Quiniæ*, B. P., which is a solution (saturated at 50° or 60° F.) of the salt in tincture of orange-peel (eight grains in the ounce). An Ammoniated Tincture (*Tinctura Quiniæ Ammoniata*, B. P.) of similar strength is made by dissolving the sulphate in proof-spirit and adding a large excess of solution of ammonia. This tincture contains quinia itself liberated from combination by the ammonia; the former tincture contains the sulphate of quinia. Quinia wine (*Vinum Quiniæ*, B. P.) is a solution of neutral sulphate and citrate of quinia in orange wine, made by dissolving the disulphate (one grain in the ounce) in orange wine by the help of citric acid. The solid official preparation of the pure disulphate is *Pilula Quiniæ*, B. P. and U. S. P., containing three parts salt to one of confection of hips. The remaining Pharmacopœial preparation of quinia is the mixed citrates of iron, ammonium, and quinia (*Ferri et Quiniæ Citras*, B. P. and U. S. P.), the well-known scale compound. It is made by dissolving ferric hydrate, prepared from ferric sulphate, and quinia, prepared from the sulphate, in solution of citric acid, ammonia also being added; the liquid, evaporated to a syrupy consistence and dried in thin layers on glass plates, yields the usual greenish-yellow scales (*vide* p. 152).

*Quiniæ Valerianas*, U. S. P., is made by dissolving precipitated

* We do not know whether or not these sulphates are ordinary sulphates, the hydrogen of the acid going over to the quinia molecule, nor whether or not the quinia molecule is univalent or bivalent; hence we cannot say whether the *common* sulphate or the *soluble* sulphate is, in constitution, the neutral sulphate. In the above paragraph the names *disulphate*, *neutral sulphate*, *acid sulphate*, indicate nothing more than that the first sulphate contains in one molecule two atoms (chemical atoms) of quinine to one of sulphuric acid, the second one of each, and the third two of acid to one of quinine.

quinia in warm aqueous solution of valerianic acid and setting aside to crystallize.

*Citrate of Quinia* has the formula $(C_{20}H_{24}N_2O_2)_2,H_3C_6H_5O_7,5H_2O$.

## Reactions.

*First Analytical Reaction.*—To a solution of quinia or its salts in faintly acid water add fresh chlorine-water and then solution of ammonia; a green coloration (thalleiochin, or dalleiochin) is produced. According to Flückiger, in very dilute solutions of quinine bromine-vapor is more useful than chlorine for this reaction.

*Second Analytical Reaction.*—Repeat the foregoing reaction, but precede the addition of ammonia by solution of ferrocyanide of potassium; an evanescent red coloration is produced (Livonius and Vogel).

*Third Analytical Reaction.*—Quinia may be impure from the presence of the other alkaloids of cinchona-bark; the following tests will determine the point. The first is Stoddart's modification of Liebig's process.

Into a glass tube or bottle put ten grains of the suspected salt, dissolve in 10 minims of dilute sulphuric acid and 60 minims of distilled water; to this add 150 minims of *pure* ether, 3 minims of spirits of wine, and 40 minims of a solution of soda (1 part of solid hydrate to 12 of water). Agitate well and set aside for twelve hours, when, if the slightest trace of quinidia or cinchonia be present, they will be seen at the line of separation between the ether and solution of sulphate of sodium.

If only a small percentage of quinidia be present, it will appear as an oily substratum, appearing under the lens as dust, from the minuteness of its particles. Cinchonia will appear more decidedly crystalline. With a little practice the eye will easily distinguish which of the alkaloids is deposited.

*Fourth Analytical Reaction.*—This is Stoddart's chemico-microscopic test. Into an ounce of distilled water drop 10 drops of dilute sulphuric acid (British Pharmacopœia strength). To this add 14 grains (or as much as will saturate the acid) of the suspected salt. Filter through paper, and to a little of the filtered solution add a few drops of solution of sulphocyanide of potassium (180 grains in 1½ ounces of water). An immediate precipitate of the several alkaloids takes place, each of which is distinct and characteristic. If quinia, quinidia, and cinchonia be present, they will all be seen on the slide, becoming more and more distinct during the first hour. A good

plan is to place on a glass slip a drop of the solution to be tested, and to put another of the sulphocyanide by its side—the drops not being larger than good-sized pin-heads. Over both place a piece of thin glass, which will cause the drops to touch. Examine the line of junction under a half-inch object-glass, when the crystals are readily seen and recognized. By this method $\frac{1}{10000}$ of a grain of quinidia or cinchonia may be easily detected. The particles arrange themselves into the respective groups—the long slender needles of the quinia salt, the round crystalline masses of the quinidia, and the large well-formed prisms of the cinchonia salts. This reaction is sufficiently constant to enable an observer who has accustomed himself to the microscopic appearance of the three pure sulphocyanides to at once distinguish the respective salts from each other.

The sulphocyanide-of-potassium solution should be of the strength indicated. If not at hand it may quickly be made, sufficiently pure for this reaction, by the following process: Cyanide of potassium (fused), sublimed sulphur, of each 120 grains; distilled water an ounce and a half. Boil in a glass flask for fifteen minutes, filter, and make up the quantity to 1½ ounce with sufficient distilled water.

A small quantity of quinia in much cinchonia or quinidia cannot be recognized by the above reaction. Therefore, before concluding that no quinia is present in a specimen of those alkaloids, the sample should be treated with ether, filtered, the solution evaporated to dryness, and the residue, if any, examined for quinia.

*Fifth Analytical Reaction.*—Formation of Iodo-sulphate of Quinine. Dissolve sulphate of quinine in weak spirit of wine slightly acidulated with sulphuric acid, and add an alcoholic solution of iodine; a black precipitate forms. Allow the precipitate to settle, pour away the fluid, wash once or twice with alcohol and then boil with alcohol; on cooling, minute crystals separate having the optical properties of the mineral tourmaline. This iodo-sulphate is sometimes termed Herapathite, from the name of its discoverer. It is so slightly soluble in water that by its means quinine can be separated from its admixture with the other cinchona alkaloids. According to Jörgensen it has the formula $4C_{20}H_{24}N_2O_2,3H_2SO_4,2HI,I_4,xH_2O$.

*Other Characters.*—Concentrated sulphuric acid dissolves quinia with production of only a faint yellow color; salicin, with which quinia may possibly be adulterated, slowly gives, under the same circumstances, a deep red.——Concentrated

nitric acid dissolves quinia, yielding a colorles solution; on heating, the solution becomes yellowish.——Quinia and its salts, heated on platinum foil, burn entirely away.——Salicin in quinia may be detected by several other tests (*vide* "Salicin" in Index).

### *Analytical Reactions of other Cinchona-alkaloids. Cinchonine. Quinidine. Cinchonidine. Quinamine.*

Quinia is analytically distinguished from the other crystalline cinchona-alkaloids by its comparatively great solubility in ether. It is distinguished from a certain "amorphous alkaloid," and alkaloids soluble in ether, by the insolubility of its iodosulphate.

*Cinchonia or Cinchonine* ($C_{20}H_{24}N_2O$) does not give the green coloration with chlorine-water and ammonia, and its solution is not fluorescent. Sulphate of cinchonia, like sulphate of quinia, when heated fuses to a resinoid mass of a rich red color. Cinchonia and quinia are distinguished from morphia by non-solubility of the alkaloid precipitated by a fixed alkali in excess of the alkaline solution. (Strychnia is also precipitated by fixed alkalies and insoluble in excess, but this alkaloid has such strongly marked reactions of its own as to preclude confusion.)

*Quinidia or Quinidine* (the Conquinine or Conchinine of Hesse) is an isomer of quinia. Its solution resembles one of quinia in being fluorescent and in giving the green coloration with chlorine-water and ammonia. Sulphate of quinidia is far more soluble in chloroform than in either of the other alkaloid-sulphates.

Quinidia is analytically separated from the other alkaloids by solution of iodide of potassium, which precipitates the extremely slightly soluble hydriodate of quinidia.

*Cinchonidia or Cinchonidine* is an isomer of cinchonia. Cinchonidia is precipitated from admixture with cinchonia and quinidia by adding tartrate of sodium, which precipitates the almost insoluble tartrate of cinchonidia. Solutions of cinchonidia are not fluorescent and do not give thalleioquin.

*Cinchovatia or Cinchovatine* occurs in a particular variety of cinchona-bark.

*Quinicia or Quinicine* and *Cinchonicia or Cinchonicine* are alkaloids produced by the action of heat on quinia or quinidia and cinchonia or cinchonidia respectively. They, also, are isomers, Hesse says polymers, of the parent alkaloids. Both yield ordinary salts. *Quiniretin* is the name given to the brown or reddish-brown indifferent substance, into which quinine in aqueous solution is converted when exposed to much light. *Quinamia or Quinamine* ($C_{20}H_{28}N_2O_2$) is a fifth cinchona alkaloid obtained by Hesse in 1872 from the bark of *Cinchona succirubra*. Its solution is not fluorescent and does not give thalleioquin.

The more important properties of the Cinchona-alkaloids are summarized in *Pharmacographia* as follows:—

| | |
|---|---|
| *a.* *Hydrated* crystals are formed by | Quinine, Quinidine. |
| No *hydrated* crystals by | Cinchonine, Cinchonidine, Quinamine. |
| *b.* *Abundantly* soluble in ether | Quinine, Quinidine (1 in 30), Quinamine, and the amorphous alkaloids. |
| *Sparingly* soluble in ether | Cinchonidine. |
| *Almost insoluble* in ether | Cinchonine. |
| *c.* *Levogyre* solutions afforded by | Quinine, Cinchonidine. |
| *Dextrogyre* solutions afforded by | Cinchonine, Quinidine, Quinamine, and the amorphous alkaloids. |
| *d.* Thalleioquin is formed by | Quinine, Quinidine, Quinicine. |
| Thalleioquin cannot be obtained from | Cinchonine, Cinchonidine, Quinamine, Cinchonicine. |
| *e.* Fluorescence is displayed by solutions of | Quinine, Quinidine. |
| No fluorescence in solutions of pure | Cinchonine, Cinchonidine, Quinamine. |

## STRYCHNIA, OR STRYCHNINE.

Formula $C_{21}H_{22}N_2O_2$. Molecular weight 334.

*Source.*—This alkaloid exists, to the extent of 0.2 to 0.5 per cent., in Nux Vomica (*Strychnos Nux Vomica*, B. P. and U. S. P.), also (Shenstone) in minute quantity in the bark of the Nux Vomica tree (false angostura bark), and, to 1.0 or 1.5 per cent., in St. Ignatius's bean (*Strychnos Ignatia*) (*Ignatia*, U. S. P.) chiefly in combination with strychnic or igasuric acid, or, after slight fermentation when moistened, with lactic acid.

*Process.*—According to the British official process for its preparation (*Strychnia*, B. P.) the nuts, disintegrated by subjection to steam and, after drying, grinding in a coffee-mill, are exhausted with spirit, the latter removed by distillation, the extract dissolved in water, coloring and acid matters precipitated by acetate of lead, the filtered liquid evaporated to a small bulk, the strychnia precipitated by ammonia, the precipitate washed, dried, and exhausted with spirit, the spirit recovered by distillation, and the residual liquid set aside to crystallize. Crystals of strychnia having formed, the mother-liquor (which contains the brucia of the seeds) is poured away, and the crystals of strychnia washed with spirit (to remove any brucia) and recrystallized.

In the U. S. P. process the rasped Nux Vomica is exhausted by very dilute hydrochloric acid, milk of lime added to the evaporated

decoction to decompose the hydrochlorate of strychnia, the precipitated and dried mixture of strychnia and lime treated with diluted alcohol to remove brucia, and then with strong hot alcohol to dissolve out strychnia; the alcohol having been recovered by distillation, the residual impure strychnia is dissolved in very dilute sulphuric acid, the solution decolorized by animal charcoal, evaporated, and set aside to crystallize, the crystals of sulphate of strychnia (*Strychniæ Sulphas*, U. S. P.) redissolved in water, ammonia added to precipitate pure strychnia, and the latter dried.

*Properties.*—Strychnia occurs "in right square octahedrons or prisms, colorless and inodorous; sparingly soluble in water, but communicating to it its intensely bitter taste; soluble in boiling rectified spirit and in chloroform, but not in absolute alcohol or in ether."

## Reactions.

*First Analytical Reaction.*—Place a minute particle of strychnia on a white plate, and near to it a small fragment of red chromate of potassium; to each add one drop of concentrated sulphuric acid: after waiting a minute or so for the chromate to fairly tinge the acid, draw the latter, by a glass rod, over the strychnia spot; a beautiful purple color is produced, quickly fading into a yellowish-red. The following oxidizing agents may be used in the place of the chromate: puce-colored oxide of lead, fragments of black oxide of manganese, ferridcyanide of potassium, or permanganate of potassium.

This reaction is highly characteristic: a minute fragment dissolved in much dilute alcohol, or better, chloroform, and one drop of the liquid evaporated to dryness on a porcelain crucible lid or other white surface, yields a residue which immediately gives the purple color on being oxidized in the manner directed.

*Other Reactions.*—Strong sulphuric acid does not act on strychnia, even at the temperature of boiling water, a fact of which advantage is taken in separating strychnia from other organic matter for purposes of toxicological analysis.——Sulphocyanide of potassium produces, even in dilute solutions of strychnia, a white precipitate, which, under the microscope, is seen to consist of tufts of acicular crystals. ——Strong nitric acid does not color strychnia in the cold, and on heating only turns it yellow.

*The Physiological Test.*—A small frog placed in an ounce of water to which $\frac{1}{100}$ of a grain of a salt (acetate) of strychnia is added, is, in two or three hours, seized with tetanic spasms on the slightest touch, and dies shortly afterwards.

Strychnia has an intensely bitter taste. Cold water dissolves only $\frac{1}{2500}$ part; yet this solution, even when largely diluted, is distinctly bitter. Alcohol is a somewhat better solvent. The salts of the alkaloid are more soluble. The official solution (*Liquor Strychniæ*, B. P.)

contains four grains of strychnia to the ounce, the solvent being three parts water, one part spirit, and a few minims (6 per ounce) of hydrochloric acid (rather more than sufficient to form hydrochlorate of strychnia).

BRUCIA, or BRUCINE ($C_{23}H_{26}N_2O_4,4H_2O$), is an alkaloid accompanying strychnia in Nux Vomica and St. Ignatius's bean to the extent of about 0.5 per cent. It is readily distinguished by the intense red color produced when nitric acid is added to it. *Igasuria* is another alkaloid of Nux Vomica. This alkaloid is said to occur in nine varieties, each slightly differing from the other in chemical composition. They resemble strychnia in physical properties.

*Curarine* ($C_{10}H_{15}N$), the active principle of the arrow-poison termed *urari*, *ourari*, *wourali*, or *woorara*, prepared from a *strychnos*, resembles strychnia in giving color by oxidation, but the color is more stable. Iodide or platino-cyanide of potassium does not with curarine afford precipitates which crystallize from alcohol like those of strychnia. Curarine, also, is soluble in water. Unlike strychnia curarine is reddened by sulphuric acid; it, also, is not dissolved out by ether from an acid or alkaline liquid.

*Distinction of Brucia from Morphia.*—The red coloration produced by the action of nitric acid on brucia is distinguished from that yielded by morphia by the action of reducing agents (such as stannous chloride, hyposulphite of sodium, sulphydrate of sodium), which decolorize the morphia-red, but change that of the brucia to violet and green (Cotton).

*Distinction of Free Alkaloids or their Salts from each Other.*—This is accomplished by remembering the appearance and other physical characters of the substances as met with in pharmacy, the effect of heat, the action of such solvents as water, alcohol, and ether, the influence of strong and diluted acids, strong and weak alkalies, oxidizing agents, chlorine-water and ammonia, and other reagents.

---

## QUESTIONS AND EXERCISES.

680. What alkaloids are more or less characteristic of the different varieties of cinchona-bark? In what form do they occur?

681. By what method is Disulphate of Quinia obtained?

682. Give the characters of disulphate of quinia.

683. Describe the tests for quinia.

684. How is the adulteration of disulphate of quinia by salicin detected?

685. Show how the sulphates of quinia or cinchonia may be proved to be present in commercial quinia.

686. How are cinchonia and quinia distinguished from morphia?

687. Whence is strychnia obtained?

688. Describe the official process for the isolation of strychnia.

689. Give the characters of strychnia.

690. Enumerate the tests for strychnia, and describe their mode of application.

691. By what reagent is brucia distinguished from strychnia?

692. Distinguish between brucia and morphia.

693. By what general methods would you distinguish common alkaloids from each other?

---

*Analytical Exercises.* — Analyze small quantities of alkaloids, their salts, and various "scale" compounds by aid of the annexed *Tables.*

---

## ALKALOIDS OF LESS FREQUENT OCCURRENCE.

ACONITIA, ACONITINA, or ACONITINE ($C_{64}H_{40}NO_2$) is an alkaloid obtained from aconite (*Aconitum Napellus*) leaves (*Aconiti Folia*, B. P. and U. S.) and root (*Aconiti Radix*, B. P. and U. S. P.). The alkaloid itself is only slightly soluble in water; it occurs in the plant in combination with a vegetable acid, forming a soluble salt.

*Process.*—The official process for its preparation (*Aconitia*, B. P. and U. S. P.) consists in dissolving out the natural salt of the alkaloid from the root by rectified spirit, recovering the latter by distillation, mixing the residue with water, filtering, precipitating the aconitia by ammonia, drying the precipitate and digesting it in ether (in which some of the accompanying impurities are insoluble), recovering the ether by distillation, dissolving the dry residue in the retort in water acidulated by sulphuric acid, again precipitating the alkaloid by ammonia, and finally washing and drying.

*Properties.*—Aconitia usually occurs as a white powder, but has been obtained and studied in the crystalline state by Groves and by Duquesnel. It is soluble in 150 parts of cold water, 50 of hot, and much more soluble in alcohol and in ether. It is one of the most violent poisons known. "When rubbed on the skin it causes a tingling sensation, followed by prolonged numbness."

The thousandth part of a grain on the tip of the tongue produces, after a minute or so, a characteristic tingling sensation and numbness: larger quantities rubbed into the skin cause numbness and loss of feeling. To a strong solution of phosphoric acid (the ordinary acid concentrated over a water-bath to a syrupy consistence) a mere trace imparts a lasting violet tint (Flückiger). Oil of vitriol turns it of a yellowish, and afterwards, dirty-violet color.

According to Wright, who, in conjunction with Groves and Williams, worked by aid of grants from the British Pharmaceutical Conference, *Aconitum napellus* yields crystalline *aconitine*, $C_{33}H_{43}NO_{12}$, crystalline *pseud-aconitine*, $C_{36}H_{49}NO_{12}$, and a non-crystalline alkaloid.

The tuberous roots of *Aconitum ferox* and other species constitute the *bish* or *bikh* of India (*Aconiti ferocis Radix*, P. I.). It chiefly contains the variety of aconitine termed *pseud-aconitine*. Much of

ALKALOI
d.D.

Make a
of $HNO_3$,
and test

*First portion.*
Add BaC
A precipit
= *Sulphu*
*Acid.*

690 DS AND THEIR SALTS, ETC.

applic

691

692

693 alkalc

| ACIDS. | SULPHURIC. HYDROCHLORIC. PHOSPHORIC. | ACETIC. MECONIC. CITRIC. |
|---|---|---|

solution with a slight excess
*An* divide it into three portions,
their as follows :—
*Table*

Make a neutral solution, divide it into two portions, and test as follows :—

| | *Second portion.* | *Third portion.* | *First portion.* | *Second portion.* |
|---|---|---|---|---|
| **A** | Add $AgNO_3$. A precipitate soluble in $NH_4HO$ = *Hydrochloric Acid.* | Boil for a time and then add a few drops to an excess of solution of molybdate of ammonium, and boil. A yellow precipitate = *Phosphoric Acid.* | Add neutral $Fe_2Cl_6$. A red color indicates acetic or meconic acid. Distinguish Decolorized by boiling = *Acetic Acid.* Not decolorized by boiling = *Meconic Acid.* | If no acetic or meconic acid, add $AgNO_3$. Precipitate (white, with a tendency to darken) soluble in $HNO_3$ = *Citric Acid.* Confirm by adding $CaCl_2$ to a neutral conc. solution and boiling, when a white precipitate of citrate of calcium separates. |

A $Cl_2$.
loid ate
B. P ic
The
plan
P
and
loid
tion,
nitia
(in w
ering
in w
by a
P
been
Duq
muc
viole
glin
T
after
ness
of f
conc
imp
yell
A
liam
fere
crys
loid
T
the
cont

[*To face page* 392.]

ALKALOIDS. { QUINIA. QUINIDIA. BEBERIA. / CINCHONIA. CINCHONIDIA. STRYCHNIA.

Dissolve a portion in $H_2O$, and add $NH_4HO$ cautiously (Preci Alkaloids (except Strychnia) and sometimes $Fe_26HO$). Add in excess (insoluble Precipitate = Cinchonia or Cinchonidia, or $Fe_26HO$). Agitate the mixture with ether and separate, by m pipette, the ethereal solution, aqueous solution, and insoluble pr

| ETHEREAL SOLUTION. | | INSOLUBLE PRECIPITATE. | AQU SOLU |
|---|---|---|---|
| May contain quinia, quinidia, or beberia. To solution in a test-tube add $H_2O$ acidulated with $HC_2H_3O_2$, and boil, burning off the ether. To a portion of the acetic solution, add Cl water and afterwards $NH_4HO$. | | Is chinchonia, cinchonidia, or $Fe_26HO$. Benzole may be employed to separate the cinchonia or cinchonidia, it being afterwards removed by heat. Dissolve in dilute $HC_2H_3O_2$, and to part of the solution add $Na_2C_4H_4O_6$, when a precipitate occurs in the case of *cinchonidia*, and no precipitate in that of *cinchonia*. The solution in $HC_2H_3O_2$ is not fluorescent, and gives no thalleioquin. | May strychn tate wit and sep chlorofo lution. rate chl solution moisten with co ed $H_2SC$ across film a s moistenly tal of en when a play of violet t lors = n nia. In doubt a $NH_4HO$ nal solu Agitat chlorofo proceed fore. |
| GREEN COLOR (thalleioquin). | NO GREEN. | | |
| Solution is fluorescent, and contains either quinia or quinidia. Concentrate the remainder of the solution and divide into two parts. To one add KI, and to the other add $(NH_4)_2C_2O_4$. KI precipitates *quinidia*, not quinia. $(NH_4)_2C_2O_4$ precipitates *quinia*, not quinidia. | To a portion of the acetic solution add KHO. A yellow precipitate = *beberia*. | | |

69 Com
appl
69
69
69
alka

A
thei
Tabl

cipitate =
$NH_4HO$
perhaps
eans of a
ecipitate.

EOUS
ITION.

$Al_2$
loid at
B. Fi
The
plan

contain
ia. Agi-
h $CHCl_3$
arate
rmic so-
Evapo-
proformic
, and
residue
ncentrat-
$)_4$. Draw
the acid
small

P
and
loid
tion,
nitia
(in
erin
in w
by a

P
beer
Duq
muc
viol
glin

ed crys-
$K_2Cr_2O_7$,
ransient
colors—
o red =
. No co-
o strych-
case of
dd
to origi-
tion.
e with
rm, and
as be-

T
after
ness
of f
conc
imp
yell

A
liam
fere
crys
loid.

T
the
cont

ıe aconitine of pharmacy is pseud-aconitine. According to Paul nd Kingzett the alkaloid of Japanese aconite has the formula $C_{29}H_{43}NO_9$.

*Unguentum Aconitiæ*, B. P., contains eight grains of the alkaloid o one ounce of prepared lard.

*Aconitum heterophyllum*, *Atis*, or *Atees* (*Aconiti heterophylli Radix*, P. I.) contains no aconitia, but an alkaloid having the formula $C_{44}H_{74}N_2O_5$ (Broughton, in *Pharmacographia*).

ATROPIA, or ATROPINE ($C_{17}H_{23}NO_3$), exists in the Belladonna, or Deadly Nightshade (*Atropa Belladonna; Belladonnæ Folia et Radix*, B. P. and U. S. P.), as soluble acid malate of atropia.

*Process.*—It is obtained in the pure state by exhausting the root with spirit, precipitating the acid and some coloring matter by lime, filtering, adding sulphuric acid to form sulphate of atropia (which is somewhat less liable to decomposition during subsequent operations than the alkaloid itself), recovering most of the spirit by distillation, adding water to the residue, and evaporating till the remaining spirit is removed; solution of carbonate of potassium is then poured in till the liquid is nearly but not quite neutral, by which resinous matter is precipitated; the latter is filtered away, excess of carbonate of potassium then added, and the liberated atropia dissolved out by shaking the liquid with chloroform. The latter solution, having subsided, is removed, the chloroform recovered by distillation, the residual atropia dissolved in warm spirit, coloring matter separated by digesting the liquid with animal charcoal, the solution filtered, evaporated, and set aside to deposit crystals.

*Solubility.*—Atropia is sparingly soluble in water, the liquid giving an alkaline reaction—more soluble in alcohol and ether.

*Tests.*—Atropia solutions give with perchloride of gold a yellow precipitate. One drop of a dilute aqueous solution (two grains to the ounce) powerfully dilates the pupil of the eye. It is generally applied on a piece of thin tissue paper or small disk placed between the eyelid and the eye. Baryta water decomposes it into *tropine* ($C_8H_{15}NO$) and *tropic acid* ($C_9H_{10}O_3$), a molecule of water being absorbed; hence the atropia so called would seem to be *tropate of atropia*.

*Preparations.*—The alkaloid itself (*Atropia*), its sulphate (*Atropiæ Sulphas*, a colorless powder soluble in water, made by neutralizing atropia with sulphuric acid), their solutions (*Liquor Atropiæ*, four grains per ounce, and *Liquor Atropiæ Sulphatis*, four grains per ounce), and an ointment (*Unguentum Atropiæ*, eight grains per ounce) are the preparations official in the British Pharmacopœia. The alkaloid and its sulphate are official in the United States Pharmacopœia.

*Dataria* or *Daturine*, an alkaloid in *Datura Stramonium*, or Thorn-apple (*Stramonii Folia et Semina*, B. P. and U. S. P.), and in *Dhatura*, *Datura alba* (*Datura Folia Semina*, P. I.); it appears to be either identical with atropia or, more likely, closely allied.

BEBERIA, BEBERINE, BEBIRINE, or BIBIRINA ($C_{19}H_{21}NO_3$), is an alkaloid existing in the bark of Bebeeru or Bebiæ (*Nectandra Rodiæi*).

*Process.*—According to the British Parmacopœia, it, or rather its sulphate, $C_{38}H_{42}N_2O_6,H_2SO_4$ (*Beberiæ Sulphas*, B. P.), may be prepared by exhausting the bark (*Nectandræ Cortex*, B. P. and U. S. P.) with water acidulated by sulphuric acid, concentrating, removing most of the acid by lime, filtering, precipitating the alkaloid by ammonia, filtering, drying, dissolving in spirit (in which some accompanying matters are insoluble), recovering most of the spirit by distillation, neutralizing by dilute sulphuric acid, evaporating to dryness, dissolving the residual sulphate in water, evaporating to the consistence of a syrup, and spreading on glass plates, drying the product at 140° F. Thus obtained, it occurs in thin dark brown translucent scales, yellow when powdered, strongly bitter, soluble in water and in alcohol.

*Tests.*—Alkalies give a pale yellow precipitate of beberia when added to an aqueous solution of a salt of the alkaloid; the precipitate is soluble in ether. With red chromate of potassium and sulphuric acid beberia gives a black resin, and with nitric acid a yellow resin. *Buxine*, from the bark of *Buxus sempervirens; Pelosine*, or *Cissampeline*, from the root (*Pareiræ Radix*, B. P.) of *Cissampelos Pareira;* and *Paricine*, from a false Para cinchona-bark, are probably identical with Beberine (Flückiger).

*Nectandria* ($C_{20}H_{23}NO_4$).—Drs. Maclagan and Gamgee have recently discovered this second alkaloid in Bebeeru bark. It differs from beberia in fusing when placed in boiling water, in being much less soluble in ether, in giving with strong sulphuric acid and black oxide of manganese a beautiful green and then violet coloration, and in having a distinct molecular weight. They are of opinion that two other alkaloids exist in Bebeeru bark.

Berberia, or Berberine ($C_{20}H_{17}NO_4$), is an alkaloid existing in several plants of the natural order *Berberideæ* (three species yield *Indian Barberry*, *Berberis Cortex*, P. I.), in Calumba-root (*Calumbæ Radix*, B. P. and U. S. P.), *Goldthread* (*Coptis trifolia*, U. S. P.) according to Mayer, in the root of *Coptis Suta*, or *Nieshnic Bitters* (*Captidis Radix*, P. I.), an Indian tonic, and in many yellow woods. *Hydrastis canadensis*, *Yellow-root* or *Golden Seal*, contains berberia, though another alkaloid, *hydrastia*, and even a third are said to be present. *Xanthorrhiza apiifolia*, an old American tonic, and, apparently, *Xanthoxylon fraxineum*, or Prickly Ash, also contain berberia. The rhizome of *Menispermum canadense*, *Yellow Parilla*, contains, according to Maisch, a colorless alkaloid as well as berberia. The color of the tissues of these vegetables is apparently due to berberia; for the alkaloid itself is remarkable for its beautiful yellow color.

*Tests.*—When a dilute solution of iodine in iodide of potassium is added to a solution of any salt of berberia in hot spirit, excess of iodine being carefully avoided, brilliant green spangles are deposited. The reaction is sufficiently delicate to form, according to Perrins, an excellent test of the presence of berberia. This iodo-compound polarizes light, and has other analogies with a similar quinine-salt termed herapathite.

Berberia is not an official alkaloid; but the plants in which it occurs are used as medicinal agents in all parts of the world.

*Process.*—Berberia is readily extracted by boiling the raw material with water, evaporating the strained liquid to a soft extract, digesting the residue in alcohol, recovering the alcohol by distillation, boiling the residue with diluted sulphuric acid, filtering, and setting aside; the sulphate of berberia separates out, and may be purified by recrystallization from hot water. The alkaloid itself is obtained by shaking hydrate of lead with a hot aqueous solution of the sulphate of berberia (Procter).

*Podophyllum-root* (*Podophylli Radix*, B. P., *Podophyllum*, U. S. P.) is said to contain berberia. In preparing the resin of podophyllum or May-apple (*Podophylli Resina*, B. P. and U. S. P.) an alcoholic extract of the root is poured into water acidulated by hydrochloric acid, which causes aggregation of the resin particles and consequent facility of extraction not otherwise easily attained. Any hydrochlorate of berberia, which is almost insoluble in dilute mineral acids, is probably precipitated with the resin (Maisch). Water is said to extract hydrochlorate of berberia from the resin. The resin is the chief active principle of podophyllum root. No acid is ordered in U. S. P.

CAPSICIA, or CAPSICINE.—M. Felletár obtained from capsicum-fruits (*Capsici Fructus*, B. P.) a volatile alkaloid having the smell of conia. Thresh has obtained crystalline hydrochlorate and sulphate. Thresh has also succeeded in isolating the active principle of capsicum, which he has termed *capsaicin* ($C_9H_{14}O_2$), a crystalline, non-alkaloidal, excessively acrid substance. Its exact chemical character is not yet made out. (See also Capsicin in Index.)

COCAINA or COCAINE ($C_{16}H_{19}NO_4$) is an alkaloid of *Erythroxylon coca*, the leaves of which are said to be a powerful restorative to the human system.

COLCHICIA, or COLCHICINE, the active principle of *Colchicum autumnale* (*Colchici Cormus; Colchici Semina*, B. P.), is said to be an alkaloid, though some investigators think it has more of the characters of a neutral substance and give it the name *Colchicin*. It needs further examination. The most active medicinal preparation is an extract made from the *fresh* seeds by digestion in large volumes of alcohol of at least 90 per cent., and subsequent digestion of the marc in hot water. The extracts left on evaporating the two fluids separately are to be carefully mixed (Mols.).

CONIA, CONYLIA, CONINE, CONICINE, or CICUTINE. Formula $C_8H_{15}N$, or $(C_8H_{14})''HN$.—This alkaloid is a volatile liquid, occurring in hemlock (*Conium maculatum*), in combination with an acid (malic ?). It is not official. According to Petit, its boiling-point is 170° C. and its density 0.846.

*Process.*—It may be obtained by distilling hemlock-fruit (*Conii Fructus*, B. P.) with water rendered slightly alkaline by caustic soda or potash, or by similarly treating the fresh juice of the leaves. The alkaloid is a yellow oily liquid, floating on the water that distils over; by redistillation it is obtained colorless and transparent.

*The salts of conia* have no odor, but when moistened with solution of an alkali yield the alkaloid, the strong smell of which, at once recalling hemlock, is characteristic. Extract of hemlock leaves (*Conii*

*Folia*, B. P. and U. S. P.), to which solution of potash and boiling water have been added, forms the official Inhalation of Conia (*Vapor Coniæ*, B. P.).

*Tests.*—Sulphuric acid turns conia purplish-red, changing to olive-green, nitric acid a blood-red; perchloride of gold produces a yellowish-white precipitate, perchloride of platinum no precipitate, in aqueous solutions.

Hemlock also contains *methyl-conia* ($C_8H_{14}$)"$CH_3N$ (Kekulé and Van Planta), and *conhydrine*, $NC_8H_{14}H_2HO$.

According to Schiff, conia, isomeric, at least, with the natural alkaloid, may be produced artificially by action of ammonia on butyric aldehyd and destructive distillation of the resulting compound.

CORYDALIA, CORYDALINA, or CORYDALINE, is an alkaloid obtained by Wenzell from Turkey Corn, the tubers of *Dicentra* (*corydalis*) *formosa*.

DATURIA, or DATURINE, *vide* ATROPIA.

DELPHIA, DELPHINE, or DELPHINIE ($C_{34}H_{35}NO_9$), the poisonous alkaloid of Stavesacre (*Delphinium staphisagria*). The powdered seeds of the plant are employed to kill the *pediculi* of animals.

DITAMINE (Jobst and Hesse), the *Ditain* of Gruppe, is the alkaloid of "Dita," or bark of *Echites scholaris* or *Alstonia scholaris* (*Alstoniæ cortex*, P. I.), a reputed febrifuge.

DUBOISIA, or DUBOISINE, is the name suggested by Gerrard for the alkaloidal active principle he extracted from *Duboisia myoporoides*, unless, as he adds, the body should prove to be atropine, to which it is at all events closely allied.

EMETIA, or EMETINE ($C_{30}H_{44}N_2O_8$; Glenard).—This alkaloid is the active emetic principle of *Cephælis ipecacuanha* (*Ipecacuanha*, B. P. and U. S. P.). It occurs in combination with ipecacuanhic acid. The nitrate is peculiarly slightly soluble in water (Lefort). In the *Pulvis Ipecacuanhæ*. B. P. and U. S. P., or "Dover's Powder" (Powdered Ipecacuanha, 1 part; Powdered Opium, 1 part; and Sulphate of Potassium, 8 parts), minute division of the active ingredients is promoted by prolonged trituration with sulphate of potassium, which is a very hard salt. The Indian substitute of Ipecacuanha is the dried leaf (*Tylophoræ Folia*, P. I.) of *Tylophora asthmatica*. Its active principle has not been satisfactorily isolated.

*Gelseminia*, or *Gelsemine* ($C_{11}H_{19}NO_2$), is the alkaloid of *Gelsemium sempervirens*, or *Carolina Yellow Jasmine* (*Gelsemium*, U. S. P.), in the tissues of which plant *æsculin* ($C_{30}H_{24}O_{19}$; the *gelseminic acid* of Wormley, the fluorescent glucoside of the Horse Chestnut, and of many other plants) is also present.

HYOSCYAMIA, or HYOSCYAMINE ($C_{18}H_{23}N_2O_3$; other authors, $C_{16}H_{22}NO_3$), a volatile alkaloid, is said to occur in the leaves (*Hyoscyami Folia*, B. P. and U. S. P.), seed (*Hyoscyami Semen*, U. S. P.), and other parts of Henbane. Its effect on the eye is similar to that of atropia. The researches of Buckheim point to the conclusion that hyoscyamine is the tropate of an alkaloid homologous with tropine (Sonnenschein).

JERVIA, or JERVINE ($C_{30}H_{46}N_2O_3$), occurs in *Veratrum album* (U.

S. P.), White Hellebore, and *V. viride* (U. S. P.),* American White Hellebore, the root of which is officially recognized in Great Britain (*Veratri viridis Radix*, B. P.). Its salts are much less soluble in water than those of veratria. According to Bullock, *Veratrum viride* contains still another alkaloid—*veratroidia;* and, according to Mitchell, *Veratrum album* also contains an alkaloid which he terms *veratralbia*. Tobien gives the formula of jervia as $C_{27}H_{47}N_2O_3$, and of veratroidia as $C_{31}H_{79}N_2O_{16}$ or $C_{24}H_{37}NO_7$.

LOBELIA, or LOBELINE.—A volatile fluid alkaloid first isolated from the dried flowering herb *Lobelia inflata* (*Lobelia*, B. P. and U. S. P.) by Procter. In the pure state it smells slightly of the plant, but mixed with ammonia it emits a strong and characteristic smell of the herb. With acids it forms crystalline salts.

LUPULINE is stated by Greismayer to be a liquid volatile alkaloid contained in hops (*Humulus lupulus*).

NECTANDRIA, *vide* BEBERIA.

NICOTIA, NICOTINA, NICOTYLIA, or NICOTINE. — Formula $C_{10}H_{14}N_2$, or $(C_5H_7)'''_2N_2$. This is also a volatile liquid alkaloid, forming the active principle of tobacco (*Nicotiana tabacum*), malate and citrate of nicotina being the forms in which it occurs in the leaf (*Tabaci Folia*, B. P. and U. S. P.). Its odor is characteristic; like conia, it yields a precipitate with perchloride of gold; but, unlike that alkaloid, its aqueous solutions are precipitated yellowish-white by perchloride of platinum. It is not official.

PHYSOSTIGMA, or PHYSOSTIGMINE ($C_{30}H_{21}N_3O_4$).—An alkaloid contained in the Calabar bean (*Physostigmatis Faba*), the seed of *Physostigma venenosum* (Jobst and Hesse). A trace of it powerfully contracts the pupil of the eye; a small quantity is highly poisonous. Fraser also isolated another (but, possibly, the same) principle, and termed it *Eseria*, from Eserĕ, the name of this ordeal-poison at Calabar.

PILOCARPINE is, apparently, the active principle of the diaphoretic and sialogogue *Jaborandi* (*Pilocarpus pinnatifolius*). The occurrence of an alkaloid in this plant was first announced by Hardy, followed almost immediately by Byasson. A crystalline nitrate and hydrochlorate were first obtained by Gerrard. Kingzett gives the formula of pilocarpine as $C_{23}H_{34}N_4O_4$. The leaves also yield an essential oil, a terpene $C_{10}H_{16}$ (Hardy).

PIPERIA, or PIPERINE ($C_{17}H_{19}NO_3$) is a feeble alkaloid occurring in *white*, *black* (*Piper Nigrum*, B. P. and U. S. P.), *long pepper* (Chavica officinarum, Mign.) and *cubeb pepper* (*Cubeba*, B. P. and U. S. P.), associated with volatile oil and resin; to these three substances the odor, flavor, and acridity belong. Piperia is obtained on boiling white pepper with alcohol, and evaporating the liquid with solution of potash, which retains resin. Recrystallized from alcohol, piperia forms colorless prisms fusible at 212°. With acids and certain metallic compounds it forms salts, and distilled with strong alkali

* "The name *Green Hellebore* is sometimes applied to the drug, but it properly belongs to *Helleborus viridis* (p. 419), which is medicinal in some parts of Europe."—*Hanbury*.

yields *piperidia* or *piperidine* ($C_5H_{10}HN$), an alkaloid of strong chemical properties, and *piperic acid* ($C_{12}H_{10}O_4$). According to Buckheim the amorphous resin of the peppers is similar in constitution to piperine, alkalies breaking it up into piperidine and *chavicic acid*. *Pyrethrin* is also said to be a member of the series. The piperine of cubeb pepper is not to be confounded with *cubebin*, a neutral constituent having the formula $C_{23}H_{24}O_{10}$.

*Sanguinarina*, or *Sanguinarine* ($C_{27}H_{29}N_4O_8$), is a colorless alkaloid obtained from the rhizome of *Sanguinaria Canadensis* (U. S. P.) or Blood-root. Its salts have a red, crimson, or scarlet color.

SOLANIA, or SOLANINE ($C_{43}H_{79}NO_{16}$).—An alkaloid said to exist in the Woody Nightshade or Bitter-sweet (*Solanum dulcamara*). The dried young branches of the plant are official (*Dulcamara*, B. P. and U. S. P.). The alkaloid is only slightly soluble in water, alcohol, or ether; nitric acid colors it yellow; sulphuric acid produces at first a yellow, then a violet, and finally a brown coloration. It is said to be a conjugated compound of sugar with *solanidine* ($C_{25}H_{39}NO$). Geissler finds *dulcamarin* ($C_{22}H_{34}O_{10}$), a glucoside, to be the bitter constituent of *Solanum dulcamara*.

SPARTEIA, or SPARTEINE ($C_{15}H_{26}N$), is a poisonous volatile alkaloid occurring in Broom-tops (*Scoparii Cacumina*, B. P. and U. S. P.). Its discoverer, Stenhouse, considers that the diuretic principle of broom is *Scoparin*, a non-poisonous body. Mills has obtained ethyl-sparteine ($C_{15}H_{25}C_2H_5N$) and diethyl-sparteine ($C_{15}H_{24}C_2H_5C_2H_5N$).

THEIA, THEINE, or CAFFEINE ($C_8H_{10}N_4O_2+H_2O$). This alkaloid occurs in tea, 1 to 4 per cent., coffee (*Caffea*, U. S. P.), .5 to 2 per cent., Maté or Paraguay tea, .2 to 2 per cent., guarana, 5 per cent., and the kola-nut. Infusions and preparations of these vegetable products are used chiefly as beverages by three-fourths of the human race. It is remarkable that the instinct of man, even in his savage state, should have led him to select, as the bases of common beverages, just the four or five plants which out of many thousands are the only ones, so far as we know, containing theia. Theia is volatile. Considerable quantities may be collected by condensing the vapors evolved during the roasting of coffee on the large scale. The infusion of tea, from which astringent and coloring-matters have been precipitated by solution of subacetate of lead, and which has been evaporated to a small bulk, yields a precipitate of theia on the addition of a strong solution of carbonate of potassium. It may be crystallized from alcohol or by sublimation.

*Test.*—Concentrated nitric acid, or a mixture of chlorate of potassium and hydrochloric acid, rapidly oxidizes theia, forming compounds which with ammonia yield a beautiful purple-red color, resembling the murexid obtained under similar circumstances from uric acid; the oxidation must not be carried too far. Theine boiled with caustic potash yields methylamine ($CH_3HHN$), the vapor of which has a peculiar, characteristic odor.

*The chemical action of theia* on the system is not yet made out. Liebig thought it might aid in the production of a substance a normal amount of which is so necessary, an abnormal so unpleasant—namely,

bile. Most chemists agree that it arrests the rapid consumption of tissue, and consequent feeling of fatigue, which is especially experienced after hard work with mind or body.

VERATRIA, or VERATRINE ($C_{32}H_{50}NO_9$, Schmidt and Koppen; $C_{52}H_{86}N_2O_{15}$, Weigelin).—This alkaloid occurs in *Cevadilla* (*Sabadilla*, B. P.; the seeds of *Asagrea officinalis*, Lindley, termed *Sabadilla officinarum* by Brandt, and *Veratrum officinale* by Schlecht). According to Weigelin, Cevadilla contains two isomeric varieties of veratria, the one soluble, the other insoluble in water. There are also present *Sabadilline* ($C_{41}H_{66}N_2O_{13}$) and *Sabatrine* ($C_{51}H_{86}N_2O_{17}$). The veratrine of trade contains the two latter alkaloids (Weigelin). A mere trace of veratria brought into contact with the mucous membrane of the nose causes violent fits of sneezing. These alkaloids and those from the different species of Veratrum are evidently very closely allied. Wright and Luff, by the use of tartaric acid, a solvent less likely than the strong acids to decompose alkaloids, extract from Cevadilla, Veratrine, $C_{37}H_{53}NO_{11}$; Cevadine, $C_{32}H_{49}NO_9$; and Cevadilline, $C_{34}H_{53}NO_8$.

*The official process* for the preparation of the alkaloid (*Veratria*, B. P.) consists in exhausting the disintegrated cevadilla seeds by alcohol, recovering most of the spirit by distillation, pouring the residue into water, by which much resin is precipitated, filtering, and precipitating the veratria from the aqueous solution by ammonia. It is purified by washing with water, solution in dilute hydrochloric acid, decolorization of the liquid by animal charcoal, reprecipitation by ammonia, washing, and drying. The U. S. P. process is similar, but includes treatment of the first crude veratria by diluted sulphuric acid and precipitation of alkaloid by magnesia.

*Unguentum Veratriæ*, B. P., contains eight grains of the slightly impure alkaloid obtained as just described, rubbed down with half a drachm of olive oil and diffused through one ounce of prepared lard.

---

## QUESTIONS AND EXERCISES.

694. How is Aconite prepared?
695. Give the strength of the official preparations of Atropia.
696. Describe the properties of atropia.
697. What is the active principle of stramonium?
698. Mention pharmacopœial substances containing beberia and berberia respectively.
699. Give the characters of beberia.
700. In what does neetandria differ from beberia?
701. Mention the characteristics of conia.
702. What is the active principle of Ipecacuanha?
703. Name the alkaloid of Tobacco.
704. Give the name and properties of the active principle of Calabar Bean.
705. What are the sources of piperia?

706. Whence is theia obtained?
707. Describe the preparations of veratria.
708. State the properties of veratria.

---

## BITTER OR TONIC SUBSTANCES, Etc.

The following articles, employed medicinally in such forms as Decoction, Extract, Infusion, Tincture, etc., contain active principles which have not yet been thoroughly examined. Some of these principles have been isolated, and a few have been obtained in the crystalline condition; but their constitution has not been sufficiently well made out to admit of the classification of the bodies either among alkaloids, glucosides, acids, or other well-marked principles.

*Andrographis Caules et Radix*, P. I., from *Andrographis paniculata*, Kariyat.
*Anthemidis flores.*
*Asclepias tuberosa.* Pleurisy root (Asclepedin).
*Aurantii Cortex* (Hesperidin).
*Azadirachtæ Cortex et Folia*, P. I., from *Azadirachta indica*, Nim or Margosa.
*Baptisia tinctoria.* Wild Indigo. Alkaloid? B. L. Smedley.
*Bonducella semina*, P. I., from *Cæsalpinia (guilandina) bonducella.* Bondue seeds or nickar nuts.
*Buchu folia.*
*Calotropis Cortex*, P. I., from *Calotropis procera* and *C. gigantea*, Mudar.
*Canellæ albæ cortex.*
*Cascarillæ cortex* (Cascarillin, $C_{12}H_{11}O_4$).
*Caulophyllum tralictroides.* Blue cohosh. Alkaloid?
*Cimicifuga (Actea)* racemosa (Cimicifugin?).
*Cusparieæ cortex.* Angostura bark (Cusparin).
*Cypripedium pubescens* (Cypripedin?).
*Euonymus Europæus.* Wahoo bark (Euonymin?).
*Gulancha (Tinosporæ radix et caules*, P. I.).
*Gynocardiæ semina*, from *Gynocardia odorata (Chaulmugra)*. P. I.
*Hydrocotyles folia*, P. I., from *Hydrocotyle asiatica*, Indian pennywort.
*Iris versicolor.* Blue flag (Iridin or Irisin?).
*Lactuca* (Lactucin).
*Lupulus.*
*Maticæ folia.*
*Quassiæ lignum* (Quassin, $C_{10}H_{12}O_3$).
*Serpentariæ radix.*
*Soymidæ Cortex*, P. I., Rohun Bark, from *Soymida febrifuga.*
*Taraxaci radix* (Taraxacin).
*Toddaliæ radix*, P. I.
*Veronica virginica* (Leptandrin?).

## AMYLACEOUS AND SACCHARINE SUBSTANCES.

### STARCH.

Formula $C_6H_{10}O_5$.

*Processes.*—Rasp or grate, or, with a knife, scrape a portion of a clean raw potato, letting the pulp fall on to a piece of muslin placed over a small dish or test-glass, and then pour a slow stream of water over the pulp; minute particles or *granules* of starch pass through the muslin and sink to the bottom of the vessel, fibrous matter remaining on the sieve. This is potato-starch. Even diseased potatoes furnish good starch by this method. Wheat-starch (*Amylum*, B. P. and U. S. P.) may be obtained by tying up some flour in a piece of calico and kneading the bag in a slow stream of water flowing from a tap, the washings running into a deep vessel, at the bottom of which the white starch collects; the sticky matter remaining in the bag is *gluten*. The *blue starch* of the shops is artificially colored with smalt or indigo, to neutralize the yellow tint of recently washed linen; it should not be used for medicinal purposes. Starch dried in mass splits up into curious columnar masses, resembling the basaltic pillars of Fingal's Cave in Staffa, or those of the Giant's Causeway in the North of Ireland. The cause of the phenomenon, which may also be seen in grain-tin, is not conclusively known.

*Gluten* is the body which gives tenacity to dough and bread. It seems to be a mixture of vegetable fibrin, vegetable casein, and an albuminous matter termed gluten. These substances and gluten itself are closely allied; each contains about 16 per cent. of nitrogen. *Wheaten Flour* (*Farina Tritici*, B. P.) contains about 72 per cent. of starch and 11 of gluten, as well as sugar, gum, fine bran, water, and ash. The compactness of barley, well seen in Husked or Pearl Barley (*Hordeum Decorticatum*, B. P. and U. S. P.), is said to be due to the large amount of vegetable fibrin present. During germination the fibrin is destroyed, hence, probably, the cretaceous character of malt. *Oatmeal* (*Avenæ Farina*, U. S. P.) is very rich in albuminoid or flesh-forming constitutents, containing nearly 16 per cent. *Sago*, U. S. P., is granulated starch from the Sago Palm. *Tapioca*, U. S. P., is granulated starch from the Bitter Casava. The white translucent grains known as Rice are the husked seeds of *Oryza sativa*. Rice (*Oryza*) and the Flour of Rice, or Ground Rice (*Oryza Farina*), are official in the Pharmacopœia of India. Rice is quite a staple article of food in tropical countries. Ground rice resembles flour of wheat in composition, but contains from 85 to 90 per cent. of starch.

*Microscopical Examination of Starches.*—All kinds of starch afford the blue color with iodine, showing their chemical similarity.

Physically, however, the granules of different starches differ from each other; hence a careful microscopical examination of any starch, or of any powder or vegetable tissue containing starch, enables the observer to state, with a high degree of probability, the source of the starch, either at once if he has much experience, or after comparing the granules in question with authentic specimens. A glance at the accompanying eight engravings* (Figs. 41 to 48) of common starches will show to what extent different starch-granules naturally differ in size, shape, general appearance, distinctness and character of the rugæ, and position of the more or less central point or hilum. While from different starches individual granules may be picked out which much resemble each other, the appearance of each starch as a whole is fairly characteristic. That is to say, each *group* of granules differs in one, two, or several characters from similar *groups* of granules of other starches.

A quarter-inch object-glass will commonly suffice for the microscopical examination of starch. A very little of the starch is mixed on a glass slide with a drop of water, a piece of thin covering-glass placed on the drop and gently pressed, so as to provide a very thin layer for observation. Instead of water diluted spirit of wine, diluted glycerine, turpentine, or other essential oil, Canada balsam, and other fluids may be used in cases where the markings or other appearances are not well defined. The illumination also of the granules may be varied, the light being reflected or transmitted, concentrated or diffused, white or colored, polarized or plain. Polarized light is especially valuable in developing differences, and in intensifying the effects of obscure markings. By polarized light the granules of potato-starch appear as if traversed by a black cross; wheat-starch granules and many others also peculiarly and characteristically influence polarized light. Distinctive characters will sometimes present themselves only when the granules are made to roll over in the fluid in which they have been temporarily mounted, or when the slide is gently warmed. Starches which have already been subjected to the influence of heat, partly as in sago or tapioca, or almost entirely as in bread, will of course differ in appearance from granules of the same starch before being dried, cooked, or torrefied. The characters of a starch will also somewhat vary according to the age and condition of the plant yielding it.

For plates and descriptions of the characters of other starches occurring in plants used for medicinal purposes, the reader is referred to works on Materia Medica, and to the indexes of Journals of Pharmacy, as well as to general works and magazines on microscopy. For engravings of starch-granules *in situ vide* Berg's *Anatomischer Atlas*, published by Gaertner, Berlin.

The student may place fair confidence in the accompanying illustrations, and in most of the published engravings of starch-granules;

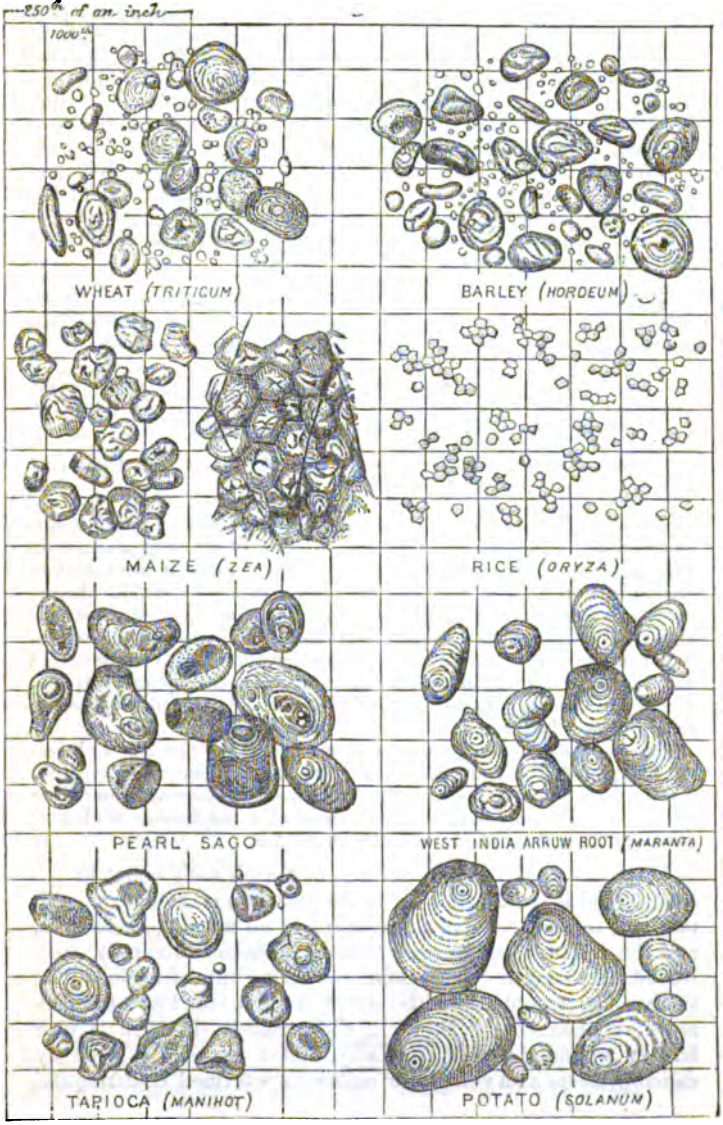
---

* By permission of Messrs. Longmans & Co., these engravings have been copied, with very few modifications, from the plates in two of the three volumes of the original edition of Pereira's "Materia Medica."

# STARCHES

*mag.ᵗ 250 diaᵐ*

*Figs. 41 to 48.*



but in microscopical analyses of importance the worker should, if possible, himself obtain actual specimens of starches for comparison from the respective seeds, fruits, and other tissues.

*Inulin* ($C_6H_{10}O_5$) is a white powder apparently occupying the place of starch in the roots of many plants, especially those of the natural order *Compositæ*. Twenty to forty-five per cent. has been obtained from elecampane (*Inula helenium*). It is also contained in the dahlia, colchicum, arnica, dandelion, chicory, etc. It is soluble in boiling water, nearly all being redeposited on cooling. Iodine turns it yellow. Long ebullition converts it into a kind of gum.

*Lichenin* ($C_6H_{10}O_5$) is a white starch-like powder largely contained in many lichens—Iceland "Moss," *Cetraria islandica*, and many others. It is soluble in boiling water, and the fluid gelatinizes on cooling. It may be precipitated from its aqueous solution by alcohol. With iodine it gives a reddish-blue color.

*Mucilage of Starch.*—Mix two or three grains of starch with first a little and then more water, and heat to the boiling-point; mucilage of starch (*Mucilago Amyli*, B. P.) results.

This mucilage or paste is not a true solution; by long boiling, however, a portion of the starch becomes dissolved. In the latter case the starch probably becomes somewhat altered.

*Chemical Test.*—To some of the mucilage add a little free iodine; a deep blue color is produced.

This reaction is a very delicate test of the presence of either iodine or starch. The starch must be in the state of mucilage; hence in testing for starch the substance supposed to contain it must be first boiled in water. The solutions used in the reactions should also be cold, or nearly so, as the blue color disappears on heating, though it is partially restored on cooling. The iodine reagent may be iodine-water or tincture of iodine. In testing for iodine its occurrence in the free state must be insured by the addition of a drop, or even less, of chlorine-water. Excess of chlorine must be avoided, or chloride of iodine will be formed, which does not color starch.

The so-called *iodide of starch* scarcely merits the name of a chemical compound, the state of union of its constituents being feeble. Substances that attack free iodine remove that element from iodide of starch. The alkalies, hydrosulphuric acid, sulphurous acid, and other reducing reagents destroy the blue color.

DEXTRIN.—Mix a grain or two of starch with about half a test-tubeful of cold water and a drop or two of sulphuric acid, and boil the mixture for a few minutes; no mucilage is formed, and the liquid, if sufficiently boiled, yields no blue color with iodine; the starch has become converted into *dextrin*. The same effect is produced if the starch is maintained at a temperature of about 320° F. for a short time. Dextrin is now largely manufactured in this way, and a paste of it used by calico-printers as a vehicle for colors; it is termed *British gum*.

The change may also be effected by *diastase*, a peculiar ferment existing in malt. Mix two equal quantities of starch with equal amounts of water, adding to one a little ground malt, then heat both slowly to the boiling-point; the mixture without malt thickens to a paste or pudding, that with malt remains thin, its starch having become converted iuto dextrin.

*Diastase* is probably the vegetable fibrin of gluten in a state of decomposition; it is so named from διάστασις (*diastasis*), *separation*, in allusion to the separation, or rather alteration, it effects among the constituent atoms of starch.

*Malt* (the word malt is said to be derived from the Welsh *mall*, soft or "rotten") is simply barley which has been softened by steeping in water, allowed to germinate slightly, and further change then arrested by the application of heat in a kiln. During germination the gluten breaks up and yields a glutinous substance termed vegetable gelatine, diastase, and other matters. To the vegetable gelatine is due much of the "body" of well-malted and slightly-hopped beer; it is precipitated by tannic acid, hence the thinness of ale (pale or bitter) brewed with a large proportion of hop or other materials containing tannic acid. A portion of the diastase reacting on the starch of the barley converts it into dextrin, and, indeed, carries conversion to the further stage of grape-sugar, as will be explained immediately. The temperature to which the malt is heated is made to vary, so that the sugar of the malt may or may not be partially altered to a dark-brown coloring material; if the temperature is high, the malt is said to be *high-dried*, and is used in porter-brewing; if low, the product is of lighter color, and is used for ale. The diastase remaining in malt is still capable of converting a large quantity of starch into dextrin and sugar; hence the makers or distillers of the various spirits operate on a mixture of malted and unmalted grain in preparing liquors for fermentation.

*Gum* is a frequent constituent of vegetable juices, existing in large quantity in several species of acacia. According to Frémy gum is a calcium salt, sometimes partially a potassium salt of the *gummic* or *arabic* radical. The formula of gummic acid is said to be $H_2C_{12}H_{18}O_{10}$, $H_2O$. Gum differs from dextrin in yielding oxalic acid, but no mucic acid when oxidized by nitric acid. *Cerasin* or *cherry-tree gum* is an insoluble modification of acacia gum. *Bassorin*, *tragan-thin* or *adraganthin* ($C_{12}H_{20}O_{10}$) is a form of gum which is insoluble in water, but absorbs a large quantity of that liquid and forms a gelatinoid mass. It occurs largely in Tragacanth, combined, like arabin, with calcium. *Pectin* or *vegetable jelly* ($C_{23}H_{40}O_{28}$, $4H_2O$) is the body which gives to expressed vegetable juices the property of gelatinizing. It forms the chief portion of Irish or Carrageen "Moss" (*Chondrus crispus*, U. S. P.). *Ceylon "Moss"* (*Gracilla-ria lichenoides* and *G. confervoides*, P. I.) contains from one-third to three-fourths of vegetable jelly.

The mucilage of marsh-mallow root (*Althea officinalis*) and of linseed or common flaxseed (*Linum usitatissimum*) is a gum-like substance containing much mineral matter. It is the basis of the

infusions termed *mallow-tea* and *linseed-tea*. Somewhat similar mucilage occurs in infusion of Bael; it is also largely yielded by the seeds of the Quince (*Cydonia vulgaris*), and by ordinary Elm-Bark (*Ulmi Cortex*, B. P.), as well as by the bark of the Red or Slippery Elm (*Cortex Ulmi Fulvæ*). *Salep*, the powdered dried tubers of many species of *Orchis*, contains a large quantity of such matter. *Squill* also. The Indian *Okra* (*Hibisci Capsulæ*, P. I., from *Hibiscus esculentus*) and *Ispaghûl* or *Spogel* seeds (*Ispaghulæ semina*, P. I., from *Plantago ispaghula*) also appear to contain a considerable quantity.

### *Isomerism. Allotropy. Polymorphism.*

The composition of dextrin is represented by the same formula as that of starch, namely $C_6H_{10}O_5$; for it has the same percentage composition as starch. Inulin (p. 404) and cellulose have also a similar formula. There are many other bodies similar in centesimal composition, but dissimilar in properties; such substances are termed *isomeric* (ἴσος, *isos*, equal, and μέρος, *meros*, part), and their condition is spoken of as one of *isomerism*. There is sometimes good reason for doubling or otherwise multiplying the formula of one of two *isomers*, *isomerides*, or isomeric bodies. Thus a molecule of olefiant gas (ethylene), the chief illuminating constituent of coal-gas, is represented by the formula $C_2H_4$, while a molecule of amylene, an anæsthetic liquid hydrocarbon, obtained from amylic alcohol, though having the same percentage composition as olefiant gas, is represented by the formula $C_5H_{10}$: for amylene, when gaseous, is about twice and a half as heavy as ethylene, and must contain, therefore, in each molecule, twice and a half as many *atoms*, for (law of Avogadro) these equal volumes must contain equal numbers of molecules; its formula is, consequently, constructed to represent those proportions. (Read again pages 38–42.) This variety of isomerism is termed *polymerism* (from πολύς *polūs*, many or much, and μέρος, part). Metastannic acid (*vide* p. 243) is a polymeric variety or *polymeride* of stannic acid. An illustration of a second variety of isomerism is seen in the case of cyanate of ammonium and urea, bodies already alluded to in connection with cyanic acid. These and several other pairs of chemical substances have dissimilar properties, yet are similar not only in elementary composition and in the centesimal proportion of the elements, but also in the fact that each molecule possesses the same number of atoms. But the reactions of these bodies indicate the probable nature of their construction; and this is shown in their formulæ by the disposition of the symbols. Thus cyanate of ammonium is represented by the formula $CNONH_4$, urea by $CON_2H_4$. Such bodies are termed *metameric* (from μετά, *meta*, a preposition denoting change, and μέρος), and their condition spoken of as one of *metamerism*. Acetate of ethyl (p. 300) is metameric with butyric acid (p. 361), for they have the same percentage composition and their vapors have the same specific gravity, and each therefore might be represented by the formula $C_4H_8O_2$; but their properties warrant us in assuming that their atoms occupy different positions in the two molecules—justify us in giving $C_2H_5C_2H_3O_2$ as a picture of a mole-

cule of acetate of ethyl, and $HC_4H_7O_2$ as a picture of a molecule of butyric acid. Acetate of methyl ($CH_3C_2H_3O_2$), propionic acid ($HC_3H_5O_2$), and formate of ethyl ($C_2H_5CHO_2$) are isomers of the metameric variety, or *metamerides;* also quinine and quinidine, cinchonine and cinchonidine, and many of the volatile oils, etc. The isomerism of starch and dextrin may be of a polymeric or of a metameric character; but we do not yet know which, and must therefore at present give them identical formulæ. Substances similar in composition and constitution, yet differing in properties, are termed *allotropic* (ἄλλος, *allos*, another, τρόπος, *tropos*, condition). Thus ordinary phosphorus, kept at a temperature of about 450° F., in an atmosphere from which air is excluded, becomes red, opaque, insoluble in liquids in which ordinary phosphorus is soluble, oxidizes extremely slowly, and only ignites when heated to near 500° F. (red or amorphous phosphorus). A black allotropic variety of phosphorus is known. There are, also, three *allotropes* of carbon which are respectively crystalline, graphitic, and amorphous. Sulphur may be obtained in the viscous as well as in the hard, brittle condition. Another illustration of *allotropy* is seen in the varieties of tartaric acid, which have different optical properties, but otherwise are identical; they are in neither of the above-mentioned states of isomerism, but are allotropic modifications of the same substance. Occasionally one and the same substance crystallizes in two distinct forms, its state is then described as one of *polymorphism* (πολύς, *polūs*, many, μορφή, *morphē*, form). Sulphur is polymorphous. It crystallizes by slow cooling in (1) prismatic crystals of sp. gr. 1.98, while in nature it occurs in (2) octahedra of sp. gr. 2.07. Melted and poured into water, sulphur takes up the form of (3) caoutchouc of sp. gr. 1.96. These differences warrant the statement that sulphur occurs in three distinct allotropic conditions. Possibly such conditions result from the association of different numbers of atoms in the molecule of the element, that is, allotropic bodies may simply be *physically polymeric* or in some other way be mere *physical isomerides.*

*Cellulin* or *cellulose*, the woody fibre of plants, familiar in the nearly pure state under the forms of "cotton-wool" (*Gossypium*, B. P. and U. S. P., "hairs of the seed of various species of Gossypium"), paper, linen, and pith, is another substance isomeric, probably polymeric, with starch. Lignin is a closely allied body, lining the interior of woody cells and vessels. By the action of nitric acid of various strengths on cellulin, peroxide of nitrogen ($NO_2$) is substituted for one, two, or three atoms of hydrogen—mono-, di-, or trinitrocellulin being formed:—

$$\underset{\text{Cellulin.}}{C_6H_{10}O_5} + \underset{\text{Nitric acid.}}{HNO_3} = \underset{\text{Mononitrocellulin.}}{C_6\left\{\begin{matrix}H_9\\NO_2\end{matrix}\right\}O_5} + \underset{\text{Water.}}{H_2O}$$

$$\underset{\text{Cellulin.}}{C_6H_{10}O_5} + \underset{\text{Nitric acid.}}{2HNO_3} = \underset{\text{Dinitrocellulin.}}{C_6\left\{\begin{matrix}H_8\\2NO_2\end{matrix}\right\}O_5} + \underset{\text{Water.}}{2H_2O}$$

$$\underset{\text{Cellulin.}}{C_6H_{10}O_5} + \underset{\text{Nitric acid.}}{3HNO_3} = \underset{\text{Trinitrocellulin.}}{C_6\left\{\begin{matrix}H_7\\3NO_2\end{matrix}\right\}O_5} + \underset{\text{Water.}}{3H_2O}$$

Trinitrocellulin is highly explosive gun-cotton; dinitrocellulin is not sufficiently explosive for use instead of gunpowder; mononitrocellulin is scarcely at all explosive. The three movable atoms of hydrogen in cellulin may be displaced by bodies other than peroxide of nitrogen.

*Dinitrocellulin* (*Pyroxylin*, B. P.) may be prepared by the following process: Mix 5 fluidounces of sulphuric acid and 5 of nitric in an earthenware mortar, immerse 1 ounce of cotton-wool in the mixture, and stir it for three minutes with a glass rod so that it is thoroughly and uniformly wetted by the acids. Transfer the cotton to a vessel containing a considerable volume of water, stir it rapidly and well with a glass rod, decant the liquid, pour more water upon the mass, agitate again, and repeat the affusion, agitation, and decantation until the washing ceases to give a precipitate with chloride of barium. Drain the product on filtering paper, and dry in a water-bath.

The *Pyroxylin*, U. S. P., is made by macerating half a troyounce of cotton for fifteen hours in a mixture of four troyounces of sulphuric acid and three and a half troyounces of nitric, and washing as above.

Pyroxylin may also be made by soaking 7 parts of white filtering paper, which has been washed in hydrochloric acid and dried, in a mixture of 140 parts of sulphuric acid (sp. gr. 1.82) and 70 of nitric acid (1.37) for three hours, and well washing the product (Guichard).

*Mononitrocellulin* and *trinitrocellulin* are insoluble in a mixture of alcohol and ether; *dinitrocellulin* or *pyroxylin* is soluble, the solution forming ordinary collodion (*Collodium*, B. P. and U. S. P.). The official proportions are 1 ounce of pyroxylin dissolved in a mixture of 36 fluidounces of ether and 12 of rectified spirit. After digesting for a few days, the liquid is decanted from any insoluble matter and preserved in a well-corked bottle. It is "a colorless, highly inflammable liquid with ethereal odor, which dries rapidly upon exposure to the air, and leaves a thin transparent film, insoluble in water or rectified spirit." Flexible collodion (*Collodium Flexile* B. P. and U. S. P.) is a mixture of collodion (6 fluidounces), Canada balsam (120 grains), and castor oil (1 fluidrachm).

---

## QUESTIONS AND EXERCISES.

709. How is wheat starch or potato-starch isolated?
710. Define gluten and glutin.
711. Enumerate the proximate principles of wheaten flour.
712. Is starch soluble in water?
713. Which is the best chemical test for starch?
714. Distinguish physically between the varieties of starch.
715. Into what compound is starch converted by heat?

716. What occurs when a mixture of starch and water is allowed to flow into hot diluted sulphuric acid?

717. If equal amounts of starch and water be heated, one containing a small quantity of ground malt, what effects ensue?

718. Write a short article on the chemistry of "malting."

719. What is the nature of gum arabic, and how is it distinguished from "British Gum"?

720. Explain *isomerism*, giving several illustrations.

721. Give examples of polymeric bodies.

722. State the formula of a body metameric with urea.

723. Define *allotropy* and *polymorphism*, giving illustrations.

724. What form of cellulin is official?

725. Mention the properties of the products of the action of nitric acids of various strengths on cellulin.

726. How is pyroxylin prepared?

---

## SUGARS.

Cane-sugar, $C_{12}H_{22}O_{11}$. Lævulose, $C_6H_{12}O_6$.
Grape-sugar, $C_6H_{12}O_6,H_2O$. Inverted-sugar, $(C_6H_{12}O_6)_2$.
Milk-sugar, $C_{12}H_{22}O_{11},H_2O$.

*Artificial formation of Grape-sugar from Cane-sugar.—Tests for Sugar.*—Dissolve a grain or two of common cane-sugar in water. To a portion of this solution placed in a test-tube add more water, two or three drops of solution of sulphate of copper, a considerable quantity of solution of potash or soda (enough to turn the color of the liquid from a light to a dark blue), and heat the mixture to the boiling-point; no obvious immediate change occurs. To another portion of the syrup add a drop of sulphuric acid, and boil for ten or twenty minutes, then add the copper solution and alkali, and heat as before; a yellowish-red precipitate of cuprous oxide ($Cu_2O$) falls. This test is exceedingly delicate.

The above reaction is due to the conversion of the *cane-sugar* ($C_{12}H_{22}O_{11}$) into *inverted-sugar*, composed of *lævulose*, $C_6H_{12}O_6$ (so called because its solution causes left-handed rotation of a ray of polarized light, cane-sugar having an opposite effect), and *grape-sugar*, $C_6H_{12}O_6,H_2O$, by the influence of the sulphuric acid, and to the reducing action of the inverted-sugar on the cupric solution. The formation of a precipitate immediately, without the action of acid, shows the presence of the latter sugars—its formation only after ebullition with acid indicating, in the absence of starch or dextrin, cane sugar. In this reduction process the sugar is oxidized and broken up into several substances; but the exact nature of the reaction has not been ascertained.

Dextrin also reduces the copper salt to suboxide, unless its solution

is cold and very dilute. It does not, however, so act on a solution of cupric acetate acidified with acetic acid, while glucose produces with this liquid the usual red cuprous precipitate (Barfoed).

*Cane-sugar* or *sucrose* (*Saccharum Purificatum*, B. P. and U.S. P.) is a frequent constituent of vegetable juices. Thus it forms the chief portion of cassia-pulp (*Cassiæ Pulpa*, B. P.), is contained in the carrot and turnip, but is most plentiful in the sugar-cane; much, however, is now obtained from the sugar-maple and beetroot. On the evaporation of the juice, common *brown* or *moist sugar* crystallizes out; this, by resolution, filtration through animal charcoal, evaporation to a strong syrup, and crystallization in moulds, yields the compact crystalline conical loaves known in trade as *lump-sugar*. From a slightly less strong syrup, slowly cooled, the crystals termed *sugar-candy* are deposited, white or colored according to the color of the syrup.

*Fruit-sugar* or *Lævulose* is uncrystallizable. It is found in the grape, fig (*Ficus*, B. P. and U. S. P.), cherry, and gooseberry; both grape-sugar and fruit-sugar in the strawberry, peach, plum, etc.

*Grape-sugar* or *glucose* (from γλυκὺς, *glucūs*, sweet) is often seen in the crystallized state, in dried grapes or raisins and other fruits; it is also the variety of sugar met with in diabetic urine. Its crystalline character is quite distinct from that of cane-sugar, the latter forming large four or six-sided rhomboidal prisms, while grape-sugar occurs in masses of small cubes or square plates. Grape-sugar is also less soluble in water but more soluble in alcohol than cane-sugar.

According to Fresenius, the percentage proportion of saccharine matter in the dried fig is 60 to 70, grape 10 to 20, cherry 11, mulberry 9, currant 6, whortleberry 6, strawberry 6, raspberry 4.

Lævulose is lævogyrate, while Sucrose and Glucose possess right-handed rotation; the latter twist a ray of polarized light from left to right, to an extent dependent on the amount of sugar present—a fact easy of application in estimating the amount of sugar in syrups or in diabetic urine.

Both cane-sugar and grape-sugar yield alcohol and carbonic acid gas by fermentation, the cane-sugar probably always passing into grape-sugar before the production of alcohol commences.

$$\underset{\text{Grape-sugar.}}{C_6H_{12}O_6} = \underset{\text{Alcohol.}}{2C_2H_5HO} + \underset{\text{Carbonic acid gas.}}{2CO_2}$$

In *bread-making*, some of the starch is converted into dextrin, and this into sugar by the ferment. The above action then goes on, the liberation of gas producing the *rising* or swelling of the mixture of flour, water, and yeast (dough)—the temperature to which the mass is subjected in the oven causing escape of most of the alcohol, and further expansion of the bubbles of carbonic acid gas in every part of the now spongy loaf. The carbonic acid gas gradually evolved when flour is worked up for bread with a mixture of dry bicarbonate of sodium and tartaric acid (best preserved by previous admixture with dried flour and a little carbonate of magnesium)—*baking-powder*—exerts similar influence. The least objectionable method of

introducing carbonic acid gas, however, is that of Danglish, whose patent Aërated Bread is made from flour by mere admixture with carbonic acid water under pressure. On removal from the cylinder, the resulting dough expands by the natural elasticity of the imprisoned carbonic acid, and the bake-oven completes the process. The crumb of bread is official (*Mica Panis*, B. P.).

*Milk-sugar* or *lactose* ($C_{12}H_{24}O_{12}$) (*Saccharum Lactis*, B. P. and U. S. P.), the sweet principle of the milk of various animals, is not susceptible of alcoholic or vinous fermentation; but it resembles grape-sugar in reducing an alkaline solution of copper with precipitation of suboxide. It is readily obtained from milk by adding a few drops of acid, stirring, setting aside for the *curds* to separate, filtering, evaporating the *whey* to a small bulk, filtering again if necessary, and allowing to cool and crystallize. It usually occurs in trade "in cylindrical masses, two inches in diameter. with a cord or stick in the axis, or in fragments of cakes—grayish-white, crystalline on the surface and in its texture, translucent, hard, scentless, faintly sweet, gritty when chewed." It is soluble in 6 parts of cold and 3 of boiling water; slightly soluble in alcohol; insoluble in ether. Powdered milk-sugar is used in pharmacy as a vehicle for potent solid medicines.

*Action of Alkali on Sugar.*—To a little solution of grape-sugar add solution of potash or soda, or solution of carbonate of potassium, and warm the mixture; the liquid is darkened in color from amber to brown, according to the amount of sugar present.

*Tests.*—The copper-reaction, the fermentation process, and the effect of alkalies form three good tests of the presence of grape-sugar, and, indirectly, of cane-sugar. A piece of merino or other woollen material, previously dipped in a solution of stannic chloride and dried, becomes of a brown or black color when dipped in a solu tion of glucose and heated to about 300° F. by holding before a fire.

*Sugar from Starch.*—Boil the starch with a little water and a drop of sulphuric acid as for dextrin, but continue the ebullition for several minutes; on testing a portion of the cooled liquid with iodine and another portion with the heated alkaline solution of a copper salt as described on page 409, it will be found that the starch has nearly all become converted into a grape-sugar, or *starch-sugar* as it is sometimes termed. When made on a large scale, a warm (131° F.) mixture of starch and water of the consistence of cream is slowly poured into a boiling solution of one part of sulphuric acid in one hundred of water, the whole boiled for some time, the acid neutralized by chalk, the mixture filtered, the liquid evaporated to a thick syrup and set aside; in a few days it crystallizes to a granular mass resembling honey. In this operation a small quantity of dextrin remains with the glucose; but if the process be con-

ducted under pressure, conversion, according to Manbré, is complete. Sugar made from the starch of rice, maize, etc., is now largely used as a partial substitute for malt in brewing. It is known as *patent sugar*, *saccharine*, dextrine-maltose, etc.

*The sugar in fresh fruits* is mainly cane-sugar; but by the action of the acid, or possibly of a ferment in the juice, it is gradually converted into *inverted sugar*, a variety differing from cane sugar in being uncrystallizable, and in having an inverted or opposite influence on polarized light, twisting the ray from right to left (lævogyrate, having lævo-rotation—hence sometimes termed *levulose*). Ripe Hips (*Rosæ Caninæ Fructus*, B. P.) contain 30 per cent. of such sugar, besides gum and acid malates and citrates. Fruit-sugar, as gathered in the form of syrup by bees, is probably a mixture of these two varieties. It is gradually altered to a crystalline or granular mass of grape-sugar, as seen in dried fruits, such as Raisins (*Uvæ*, B. P., *Uva Passa*, U. S. P.), and the Prune (*Prunum*, B. P. and U. S. P.), and in solidified honey (*Mel*, B. P. and U. S. P.). This, the common form of grape-sugar, is dextrogyrate, and hence is sometimes termed *dextrose*, to distinguish it from levulose. Honey often contains flocculent matters which cause it to ferment and yield mannite (Stoddart), alcohol, and acetic acid; hence for use in medicine it is directed (*Mel Depuratum*, B. P. and U. S. P.) to be clarified by melting and straining, while hot, through flannel previously moistened with warm water. A mixture of clarified honey 80 per cent., acetic acid 10 per cent., and water 10 per cent. is official under the name of *Oxymel* (from ὀξύς, *oxus*, acid, and μέλι, honey). A similar mixture of honey with acetic acid containing the soluble portions of squill bulbs (*Scilla*, B. P. and U. S. P.) is known as Oxymel of Squill (*Oxymel Scillæ*, B. P.). Honey or sugar-cane are the bases of the official *Confections*.

"*Honey Dew*" is a viscid saccharine matter occasionally met with on the leaves of the lime, maple, black alder, rose, and other trees. Sometimes it is sufficiently abundant to dry and fall on the ground, forming a veritable "shower of manna." It is a mixture of cane-sugar, inverted sugar, and dextrin.

*Barley sugar* is made by simply heating cane-sugar till it fuses, a change from the crystalline to the uncrystallizable condition occurring. *Treacle* (*Theriaca*, B. P.), *Molasses* (*Syrupus Fuscus*, U. S. P.), or *Melasses* (from *Mel*, honey), chiefly results from the application of too much heat in evaporating the syrups of the sugar-cane; it is a mixture of cane-sugar with uncrystallizable sugar and coloring-matter. Liquorice-root (*Glycyrrhizæ Radix*, B. P.) contains a considerable quantity of uncrystallizable sugar.

*Caramel.*—Carefully heat a grain or two of sugar in a test-tube until it blackens; the product is *caramel* or *burnt sugar* (the *Saccharum Ustum* of pharmacy). It is used as a coloring agent for gravies, confectioneries, spirits, and similar materials.

*Mannite* ($C_6H_{14}O_6$).—Boil manna with alcohol, filter, and set aside; mannite separates in colorless shining crystals or

acicular masses, to the extent of from 60 to 80 per cent. of the manna.

*Manna*, B. P. and U. S. P., is "a concrete saccharine exudation from the stem of *Fraxinus Ornus* and *F. rotundifolia;* it is obtained by making incisions in the stem of the trees." It occurs in "stalactiform pieces from one to six inches in length, and one or two inches in width, uneven, porous, and friable, curved on one side, of a yellowish-white color, with a faintly nauseous odor, and a sweetish taste." Mannite is also met with in celery, onions, asparagus, certain fungi and sea-weeds, occurs in the exudations of apple and pear trees, and is produced during the viscous fermentation of sugar.

Mannite is an alcohol, the radical of which is sexivalent $(C_6H_8)^{VI}$-6HO (Wanklyn). It is closely related to the sugars, glucose becoming mannite by action of nascent hydrogen:—

$$\underset{\text{Glucose.}}{C_6H_{12}O_6} + \underset{\text{Hydrogen.}}{H_2} = \underset{\text{Mannite.}}{C_6H_{14}O_6}.$$

Indeed, glucose itself is probably an alcohol of another radical $(C_6H_8)^{VI}6HO$. Mannite does not undergo vinous fermentation in contact with yeast. It is soluble in 5 times its weight of cold water.

*Mucic Acid* ($H_2C_6H_8O_8$) and *Saccharic Acid* ($H_2C_6H_6O_6$) are two isomeric bodies formed by the action of dilute nitric acid on sugar and mannite.

---

## QUESTIONS AND EXERCISES.

727. How are cane-sugar and grape-sugar analytically distinguished?

728. Describe the methods of extracting and purifying cane-sugar.

729. Mention the chief sources of cane-sugar.

730. Give chemical explanations of the different processes of bread-making.

731. How is milk-sugar obtained, and in what respects does it differ from other sugar?

732. By what process may starch be entirely converted into sugar?

733. What is the difference between fruit-sugar and honey?

734. What is Oxymel?

735. Describe the effect of heat on cane-sugar.

736. Describe the source and character of manna.

737. Give the latest view of the constitution of mannite.

738. Whence are mucic and saccharic acids obtained?

---

## THE GLUCOSIDES.

*Source.*—The glucosides are certain proximate vegetable principles which, by ebullition with dilute acid, or other method of decomposition, take up the elements of water and yield glucose, accompanied by a second substance, which differs in each case according to

the body operated on. Several of the glucosides which are of pharmaceutical interest will now be considered. Tannin, or tannic acid, is also a glucoside; it has been described among the acids.

*Note on Nomenclature.*—The first syllable of the names of glucosides and neutral principles generally is commonly given in allusion to origin; the last syllable is *in*, which sufficiently distinguishes them as a class.

AMYGDALIN ($C_{20}H_{27}NO_{11},3H_2O$).—This body, obtained by Robiquet, and Boutron-Charlard in 1830, was the first discovered glucoside (Liebig and Whöler, 1837). It is a white crystalline substance, existing in the bitter (*Amygdala Amara*, B. P. and U. S. P.), but not in the sweet almond (*Amygdala Dulcis*, B. P. and U. S. P.). About 2 per cent. is readily extracted by strong alcohol from the cake left when the fixed oil has been expressed from bitter almonds. From the concentrated alcoholic solution ether precipitates the amygdalin.

Make an emulsion of two or three sweet almonds by bruising and rubbing them with water, and notice that it has no odor of essential oil of bitter almonds: add a grain or two of amygdalin, an odor of essential oil of bitter almonds is at once developed. Bruise two or three bitter almonds and rub with water; the volatile oil is again developed (*Oleum Amygdalæ Amaræ* U. S. P.).

*Bitter Almond-water* (*Aqua Amygdalæ Amaræ*, U. S. P.) is made by filtering a mixture of 16 minims of the oil with 60 grains of carbonate of magnesium and 2 wine-pints of distilled water.

The source of the hydride of benzoyl, or essential oil of bitter almonds, in these reactions is the amygdalin, which, under the influence of *synaptase* or *emulsin*, a nitrogenous casein-like ferment existing in both bitter and sweet almonds, splits up into the essential oil, hydrocyanic acid, and glucose:—

$$\underset{\text{Amygdalin.}}{C_{20}H_{27}NO_{11}} + \underset{\text{Water.}}{2H_2O} = \underset{\text{Hydride of benzoyl.}}{C_7H_5OH} + \underset{\text{Hydrocyanic acid.}}{HCN} + \underset{\text{Glucose.}}{2C_6H_{12}O_6}$$

As each molecule of amygdalin yields one of hydrocyanic acid, a simple calculation shows that 17 grains (mixed with emulsion of sweet almonds) will be required to form one grain of real hydrocyanic acid, a quantity equivalent to 50 minims of the dilute hydrocyanic acid of the British Pharmacopœia.

*Test.*—The reaction between synaptase and amygdalin is applicable as a test of the presence of one by the addition of the other, even when mixed with much organic matter.

*Cherry-Laurel-water* (*Aqua Laurocerasi*, B. P., by distillation with water from *Laurocerasi Folia*, B. P.) contains hydrocyanic acid derived from a reaction similar to, indeed probably identical with, that just described, for bitter-almond oil is simultaneously pro-

duced. But the proportion of amygdalin or analogous body in cherry-laurel-leaves is most variable; hence the strength of the water is highly uncertain.

*Cortex Pruni Serotinæ*, U. S. P., the recently dried Wild Black Cherry Bark, also furnishes by distillation an essential oil and hydrocyanic acid; quince-seeds also (*Cydonia vulgaris*). The Wild Black Cherry, *Cerasus* (*Prunus*) *Virginiana*, contains amygdalin.

*Caution.*—Essential oil of almonds is of course highly poisonous. The purified oil or hydrate of benzoyl is almost innocuous; it is obtained on distilling the crude oil with milk of lime and ferrous chloride and thoroughly drying the product by shaking with fused chloride of calcium. Artificial oil of bitter almonds or nitrobenzol ($C_6H_5(NO_2)$) when taken in quantity has been known to produce death. The presence of nitrobenzol in oil of bitter almonds is detected by adding a little of the oil to a mixture of zinc and diluted sulphuric acid, shaking well, setting aside for an hour or two, filtering off the clear liquid and adding a little chlorate of potassium; a violet color (actual mauve) is produced. Or the specimen may be shaken with bisulphide of sodium to fix the essential oil, and then with ether, which dissolves out, and on evaporation will yield the nitrobenzol.

ARBUTIN ($C_{24}H_{32}O_{14}H_2O$) is contained in leaves of *Arctostaphylos uva ursi* and *Chimaphila umbellata*. It is a bitter neutral body occurring in acicular crystals, and resolvable by acids into *hydroquinone* ($C_6H_6O_2$) and glucose, and by gentle oxidation into *quinone* ($C_6H_4O_2$) and formic acid. *Ericolin* ($C_{84}H_{86}O_{21}$) is another bitter glucoside in bearberry leaves.

CATHARTIC ACID.—"The glucoside acid that now is known to confer on the Senna of Alexandria (*Senna Alexandriana*, B. P.), Tinnevely (*Senna Indica*, B. P., *Senna*, U. S. P.), and probably on American Senna (*Cassia Marilandica*, U. S. P.), its purgative property, has been named by its discoverers (Dragendorf and Kubly) Carthartic acid. Its formula has been stated as $C_{180}H_{192}N_4SO_{97}$, which, if true, accounts for its extreme stability. It is insoluble in water, strong alcohol, and ether, but enters readily into watery solution when combined with alkaline and earthy bases, in which state it exists in senna. Its ammonium salts give brownish flocculent precipitates with salts of silver, tin, mercury, copper, and lead. Antimonial salts, tannin, yellow and red prussiates, have no effect upon it. Alkalies, aided by heat, act destructively upon it. Boiled with a mineral acid it splits into a peculiar kind of glucose and an acid that has been named Cathartogenic; its formula is said to be $C_{132}H_{116}N_4SO_{44}$. The natural cathartate occurring in senna is prepared by partially precipitating by strong spirit a water infusion of senna, concentrated to a syrupy state by evaporation *in vacuo*. The filtrate is now treated with a much larger bulk of absolute alcohol, and the precipitate thus obtained is purified by repeated solution in water and precipitation by alcohol. To obtain the pure acid, advantage is taken of its colloidal properties; the crude cathartate is dissolved in moderately strong hydrochloric acid, and subjected to

dialysis on a diaphragm of parchment paper. The minimum dose of this pure acid was found to be about 1½ grain, which caused several stools with decided griping.

"The cathartic combinations that I have made are, the cathartate of ammonium, prepared from cathartate of lead by my original process, and the mixed cathartates, prepared according to Dragendorf's method as modified by myself. Of the former nearly pure salt, I have found 3¾ grains to purge fairly as to amount, but slowly as to time, and with considerable griping. Of the latter 7½ grains purged violently with much griping and sickness, which continued through the greater part of the day. It obviously would be improper to combine senna with any of its metallic precipitants, should such be desired, which is not likely. It is here satisfactory to observe that the cathartate of magnesium is soluble, and that the old-fashioned black draught agrees with new-fashioned science." (Groves.)

*Buckthorn juice* (*Rhamni Succus*, B. P.) owes its cathartic properties to a substance apparently identical with cathartic acid.

COLOCYNTHIN ($C_{56}H_{84}O_{23}$?).—This substance is the active bitter and purgative principle of colocynth-fruit (*Colocynthidis Pulpa*, B. P., *Colocynthis*, U. S. P.); it is soluble in water and alcohol, but not in ether. By ebullition with acids it furnishes glucose and a resinoid body.

CONVOLVULIN.—See JALAPIN.

COTOIN ($C_{22}H_{16}O_6$) appears to be the chief active principle of Coto bark, a Bolivian remedy for diarrhœa.

DAPHNIN ($C_{21}H_{34}O_{19}$) is the crystalline glucoside of the bark of *Daphne mezereum* (*Mezerei Cortex*, B. P.). Boiled with dilute acids it yields *daphnetin* ($C_{19}H_{14}O_9$) and glucose. The acrid principle of mezereon is resinoid.

DIGITALIN ($C_{27}H_{45}O_{15}$, Kosmann; $C_5H_8O_2$, Schmiedeberg).—This is an active principle of the Foxglove (*Digitalis*, B. P.). Boil a grain of digitalin (*Digitalinum*, B. P. and U. S. P.) with sulphuric acid for some time; flocks of *digitaliretin* ($C_{15}H_{25}O_5$) separate, and glucose may be detected in the liquid.

$$\underset{\text{Digitalin.}}{C_{27}H_{45}O_{15}} + \underset{\text{Water.}}{2H_2O} = \underset{\text{Digitaliretin.}}{C_{15}H_{25}O_5} + \underset{\text{Glucose.}}{2C_6H_{12}O_6}$$

*Properties.*—Digitalin occurs "in porous mammillated masses or small scales, white, inodorous, and intensely bitter, readily soluble in spirit, but almost insoluble in water and in pure ether, dissolves in acids, but does not form with them neutral compounds; its solution in hydrochloric acid is of a faint yellow color, but rapidly becomes green. It leaves no residue when burned with free access of air. It powerfully irritates the nostrils, and is an active poison." According to Pettenkofer an intense red color is produced if a trace of digitalin dissolved in water is mixed with a weak aqueous solution of inspissated bile and sufficient oil of vitriol added to raise

the temperature to 158° F. Moistened with sulphuric acid and the liquid exposed to the vapor of bromine, a violet color is produced.

*Process.*—The official process for the preparation of digitalin consists in dissolving the glucoside out of the digitalis leaf (*Digitalis Folia*, B. P. and U. S. P.) by alcohol, removing the alcohol by distillation, dissolving the residue in water by the help of a small quantity of acetic acid, removing much of the color from the solution by animal charcoal, neutralizing most of the acetic acid by ammonia, precipitating the digitalin by tannic acid (with which it forms an insoluble compound), washing the precipitate, rubbing and heating it with spirit and oxide of lead (which removes the acid in the form of insoluble tannate of lead), again decolorizing by animal charcoal, evaporating to dryness, washing out impurities still remaining by ether, and drying the residual digitalin. In this form digitalin is uncrystallizable.

*Pure Digitalin.*(?)—On treating commercial digitalin with chloroform only an inert substance remains undissolved. The solution yields pure digitalin on evaporation; it may be crystallized from spirit in radiating needles. (Nativelle.) The therapeutic effect of the pure substance is identical with the preparations of digitalis, but, as might be expected, more constant in its action and of course intensely powerful. *Digitoxin* ($C_{21}H_{33}O_7$) is a highly poisonous substance extracted from Foxglove by Schmiedeberg. The same chemist regards commercial digitalin from Foxglove-seeds as composed of three glucosides, namely, pure *Digitalin* ($C_5H_8O_2$), *Digitonin* ($C_{31}H_{52}O_{17}$) closely allied to saponin, and *digitalein*.

ELATERIN ($C_{20}H_{28}O_5$).—Boil elaterium (*Elaterium*, B. P. and U. S. P.), the dried sediment from the juice of the squirting-cucumber fruit (*Ecbalii Fructus*, B. P.), in a small quantity of spirit of wine, and filter; fibrous and amylaceous matters remain insoluble, while elaterin and resin are dissolved. The filtrate, concentrated and poured into a warm solution of potash, yields, on cooling, crystals of elaterin, resin being retained by the alkali. It is purified by recrystallization from spirit. Boil elaterium in dilute sulphuric acid for an hour or two, filter, and test the clear liquid for glucose; a reddish precipitate of cuprous oxide falls. This reaction is readily obtained with elaterium, but not always with elaterin; hence probably the latter is not a true glucoside. Walz states that elaterium also contains *prophetin*, a true glucoside.

Elaterin is the active principle of the so-called elaterium. Elaterium occurs "in light friable slightly incurved cakes, about one line ($\frac{1}{12}$ inch) thick, greenish-gray, acrid and bitter; fracture finely granular." Good specimens of this drug should yield, according to the British Pharmacopœia, not less than 20 per cent. of elaterin by the above process. Elaterium is sometimes adulterated with chalk and other substances.

According to the experience of Flückiger and Hanbury the best method of obtaining elaterin is to exhaust elaterium with chloroform and then to add ether to the chloroform, when crystalline elaterin is precipitated. It should be washed with a little ether and crystallized from chloroform. When pure it occurs in hexagonal scales or prisms.

*Test.*—A little is placed in a watch-glass with a drop or two of liquefied carbolic acid, and then two or three drops of strong sulphuric acid; a carmine color is developed (Lindo).

GENTIOPICRIN, or GENTIAN-BITTER ($C_{20}H_{30}O_{12}$), the neutral crystalline principle of the root *Gentiana lutea* (*Radix gentianæ*, B. P.). It is soluble in water and weak spirit. Alkalies decompose it. Dilute acids convert it into *gentiogenin* and glucose. Gentian root also contains a crystalline acid ($HC_{14}H_9O_5$) termed *gentianic* or *gentisic acid* or *gentisin*. Fused potash etc. gives with the latter an acid ($C_7H_6O_4$) which has also unfortunately been called gentisic acid.

GLYCYRRHIZIN ($C_{24}H_{36}O_9$).—Liquorice-root (*Glycyrrhizæ Radix*, B. P., and *Glycyrrhiza*, U. S. P.), in addition to uncrystallizable sugar, contains a sweet substance, *glycyrrhizin*, which, when boiled with hydrochloric acid or dilute sulphuric acid, yields a resinoid bitter body, *glycyrretin*, and an uncrystallizable sugar resembling glucose. Glycyrrhizin is only slightly soluble in cold water, hence is almost wholly excluded from the official evaporated infusion (*Extractum Glycyrrhizæ*, B. P., and *Extractum Glycyrrhizæ Liquidum*, B. P.), but is present in considerable quantity in the evaporated decoction (*Stick Liquorice*, *Spanish Liquorice*, or *Solazzi Juice*). The tropical substitute for liquorice is the root of *Abrus precatorius*, or *Indian Liquorice* (*Abri Radix*, P. I.), which also apparently contains glucose and glycyrrhizin. Glycyrrhizin has considerable power of disguising nauseous flavors. Roussin refers the sweet taste of liquorice not to pure glycyrrhizin, but to a combination of glycyrrhizin with alkalies, and states that ammoniacal glycyrrhizin has exactly the sweetness of liquorice-root.

GUAIACIN.—Resin of guaiacum (*Guaiaci Resina*, B. P. and U. S. P.), an exudation from the wood (*Guaiaci Lignum*, B. P. and U. S. P.) of *Guaiacum officinale*, is probably a mixture of several substances, among which are *Guaiaretic*, or *Guaiaretinic acid* ($C_{20}H_{26}O_4$) (Illasiwetz), *Guaiaconic acid* ($C_{22}H_{40}O_{10}$) (Hadelich), and *Guaiacin*, a glucoside. On boiling guaiacum-resin with dilute sulphuric acid for some time, glucose is found in the liquid, a green resinous substance

(*guaiaretin*) remaining insoluble (Kosmann). Most oxidizing agents, and even atmospheric air, especially under the influence of certain organic substances, produce a blue, then green, and finally a brown color, when brought into contact with an alcoholic solution of guaiacum-resin.

These effects are said to be due to three stages of oxidation (Jonas). They may be observed on adding the solution to the inner surface of a paring of raw potato.

HELLEBORIN ($C_{36}H_{48}O_6$) and HELLEBOREIN ($C_{28}H_{44}O_{15}$) are crystalline glucosides occurring in the roots of Black Hellebore (*Helleborus niger*), or Christmas Rose, and Green Hellebore (*H. viridis*), ranunculaceous herbs.

JALAPIN ($C_{31}H_{50}O_{16}$) AND CONVOLVULIN ($C_{34}H_{56}O_{16}$).—According to Keyser and Meyer, jalap-resin contains two distinct substances—convolvulin, chiefly obtained from Mexican male jalap (Ipomœa orizabensis), and jalapin, most largely contained in the true jalap (Ipomœa purga); the former is soluble in ether, the latter insoluble. Boil jalap-resin with dilute sulphuric acid for some time, and filter; a substance, which is probably a mixture of *jalapinol* ($C_{13}H_{24}O_3$) and *convolvulinol* ($C_{16}H_{30}O_3$), separates, and glucose may be detected in the clear liquid. (It is to be regretted that the authors transpose the above names, terming the old well-known jalapin convolvulin.)

$$\underset{\text{Jalapin.}}{C_{31}H_{50}O_{16}} + \underset{\text{Water.}}{5H_2O} = \underset{\text{Jalapinol.}}{C_{13}H_{24}O_3} + \underset{\text{Glucose.}}{3C_6H_{12}O_6}$$

*Jalapic Acid.*—This is contained in the portion of jalap-resin soluble in ether. It may also be obtained from jalapin by ebullition with alkalies:—

$$\underset{\text{Jalapin.}}{2C_{31}H_{50}O_{16}} + \underset{\text{Water.}}{3H_2O} = \underset{\text{Jalapic acid.}}{C_{67}H_{108}O_{35}}$$

*Jalap-resin* (*Jalapæ Resina*, B. P. and U. S. P.) is obtained by digesting and percolating jalap tubercles (*Jalapa*, B. P. and U. S. P.) with spirit of wine, adding a little water, distilling off the spirit, pouring away the aqueous portion which contains much saccharine matter, and washing and drying the residual resin. The tincture is sometimes decolorized by animal charcoal, and the evaporated product sold as jalapin.

Jalap-resin is insoluble in oil of turpentine; common resin, or rosin, soluble. If the presence of the latter is suspected, the specimen should be powdered, digested in turpentine, the mixture filtered, and the filtrate evaporated; no residue, or not more than yielded by the turpentine itself, should be obtained.

*Tampico Jalap*, from *Ipomœa simulans*, yields a resin, which apparently is chiefly convolvulin, but sometimes contains jalapin;

for a sample obtained by Hanbury was entirely soluble in ether, and another extracted by Umney was almost wholly soluble, while Evans purified some, half only of which was soluble.

The *Kaladana resin* or *Pharbitisin* of India (from *Pharbitis Nil*, P. I.) is a cathartic analogous to, if not identical with, resin of jalap.

PICROTOXIN ($C_5H_6O_2$, Oppermann; $C_{12}H_{14}O_6$, Pelletier and Couerbe) is a crystalline bitter poisonous principle (*πικρὸς*, *picros*, bitter, and *τοξικὸν*, *toxicon*, poison) occurring in *Cocculus indicus*, the dried fruits of *Anamirta cocculus*, P. I. Ludwig regarded it as a glucoside, but its constitution is not yet satisfactorily ascertained.

SALICIN ($C_{13}H_{18}O_7$).—This substance is contained in and easily extracted from willow-bark.

*Tests.*—1. To a small portion of salicin placed on a white plate or dish add a drop of strong sulphuric acid; a deep red color is produced.

2. Boil salicin with dilute sulphuric acid for some time; it is converted into *saligenin* ($C_7H_8O_2$) and glucose.

$$\underset{\text{Salicin.}}{C_{13}H_{18}O_7} + \underset{\text{Water.}}{H_2O} = \underset{\text{Saligenin.}}{C_7H_8O_2} + \underset{\text{Glucose.}}{C_6H_{12}O_6}$$

Examine a portion of the solution for grape-sugar by the copper-test.

3. To another portion of the liquid, carefully neutralized, add a persalt of iron; a purplish-blue color is produced, due to the reaction of the saligenin and the ferric salt.

4. Heat a mixture of about 1 part of salicin, 1 of red chromate of potassium, 1½ of sulphuric acid, and 20 of water in a test-tube; a fragrant characteristic odor is evolved, due to the formation of hydride of salicyl ($C_7H_5O_2H$), an essential oil identical with that existing in meadow-sweet (*Spiræa ulmaria*) and in heliotrope.

$$\underset{\text{Saligenin.}}{2C_7H_8O_2} + \underset{\text{Oxygen.}}{O_2} = \underset{\substack{\text{Hydride of}\\ \text{salicyl.}}}{2C_7H_5O_2H} + \underset{\text{Water.}}{2H_2O}$$

SANTONIN ($C_{15}H_{18}O_3$).—This substance is apparently the anhydride of a weak acid (Hesse) insoluble in ammonia, but forming a soluble calcium salt. Indeed, by boiling santonin for twelve hours with baryta-water, Cannizarro has obtained a salt from which hydrochloric acid separates *santonic acid*. From a solution of santonate of calcium the santonin is precipitated by acids. Boiled for some time with dilute sulphuric acid it yields 87 per cent. of an insoluble resinous substance (*santoniretin*) and glucose (Kosmann). Santonin (*Santoninum*, B. P. and U. S. P., and *Trochischi Santonini*, U. S. P.) is official. It is soluble in an aqueous solution of twice its weight of carbonate of sodium.

*Process.*—The process for its preparation consists in boiling *santonica*, B. P. and U. S. P. (the unexpanded flower-heads of an un-

determined species of *Artemisia*—*A. cina*, U. S. P.), with milk of lime (whereby santonate of calcium is formed), straining, precipitating the santonin or santonic acid by hydrochloric acid (acetic acid, U. S. P.), washing with ammonia to remove resin, dissolving in spirit and digesting with animal charcoal to get rid of coloring-matter, setting the spirituous solution aside to deposit crystals of santonin, and purifying by recrystallization from spirit (Mialhe).

*Test.*—To highly dilute solution of perchloride of iron add an equal bulk of concentrated sulphuric acid. Add this reagent to the santonin, or to the powder or substance suspected to be santonin, and cautiously apply heat. A red, purple, and finally violet color is produced (Linds).

SAPONIN ($C_{12}H_{20}O_7$?) is a peculiar glucoside occurring in Soapwort, the root of the common Pink, and many other plants; its solution in water, even though very dilute, froths like a solution of soap. Pereira considered *smilacin* (*Salseparin* or *parallin*), one of the principles of the supposed activity of Sarsaparilla (*Sarzæ Radix*, B. P., *Sarsaparilla*, U. S. P.), to be closely allied to, if not identical with saponin.

Saponin is also met with in the root of *Polygala Senega* (*Senegæ Radix*, B. P., *Senega*, U. S. P.), though the active principle of senega is said to reside in *polygalic acid*, probably a glucoside derivative of saponin.

According to Klunge (*Pharmacographia*), parallin, by action of acids, yields *parigenin*. The aqueous solutions of parallin froth when shaken.

SCAMMONIN ($C_{33}H_{32}O_{16}$).—Boil resin of scammony (*Scammoniæ Resina*, B. P. and U. S. P.) with dilute sulphuric acid for some time; glucose may then be detected in the liquid, a resinous acid termed *scammoniol* ($C_{14}H_{13}O_3$?) being produced at the same time.

Natural scammony (*Scammonium*, B. P.) is an exudation from incisions in the living root (*Scammoniæ Radix*, B. P.) of *Convolvulus Scammonia*. It contains from 10 to 20 per cent. of gum, and, therefore, when rubbed up with water, gives an emulsion. "Ether removes from 80 to 90 per cent. of resin" (B. P.). The official resin of scammony contains no gum, and therefore gives no emulsion when rubbed up with water. It is made by digesting the root in spirit, adding water, distilling off the alcohol, and washing the residual resin with hot water till free from gum. There seems to be little or no chemical difference between the extracted resin and the resin of the exuded scammony.

Resin of scammony is soluble in all proportions in ether. Spirgatis states that it is identical with the resin of Mexican Male Jalap, which also is soluble in ether. Sulphuric acid slowly reddens it. It is said to be liable to adulteration with resin of true jalap, guaiacum-resin, and common rosin. Resin of true jalap is insoluble in ether, guaiacum-resin is distinguished by the color-tests mentioned under GUAIACIN, and rosin by the action of sulphuric acid.

SCILLITIN.—Schroff believes the bitter principle of the squill-bulb (*Scilla*, B. P.) to be a glucoside. Squill also contains a large quantity of mucilage.

The bulbous root of *Crinum asiaticum* is official in the Pharmacopœia of India (*Crini Radix*, P. I.) as a substitute for squill. It has not been chemically investigated.

## QUESTIONS AND EXERCISES.

739. Define glucosides, and mention those of pharmaceutical interest.

740. Draw out an equation illustrative of the development of Oil of Bitter Almonds.

741. How much pure amygdalin will yield one grain of real hydrocyanic acid?

742. To what does Cherry-Laurel-water owe its activity? Is the preparation trustworthy?

743. Mention the active principle of Senna.

744. By what process is the glucoside of the purple foxglove prepared?

745. State the circumstances under which Guaiacum-Resin and Jalap-Resin yield glucose.

746. Mention a test for guaiacum-resin.

747. How may the adulteration of jalap-resin by rosin be detected?

748. Enumerate the tests for Salicin.

749. How is santonin officially prepared?

750. Name sources of saponin.

751. What is the difference between Scammony and Resin of Scammony?

752. How would you detect resins of turpentine, guaiacum, or jalap, in resin of scammony?

## ALCOHOL AND ALLIED BODIES.

### ALCOHOL, OR HYDRATE OF ETHYL.

*Formation of Alcohol.*—Ferment two or three grains of sugar by dissolving in a test-tube full of water, adding a little yeast (*Cerevisiæ Fermentum*, B. P., *Fermentum*, U. S. P.), or a piece of the so-called German or dried yeast, and setting the whole aside for several hours in a warm place at a temperature of 70° or 75°; carbonic acid gas is evolved, and, if the tube be inverted in a small dish containing water, may be collected in the upper part of the tube and subsequently tested: the solution contains alcohol. If the experiment be made on larger quantities (four ounces of sugar, one of yeast, and a pint of water) the fermented liquid should be distilled, one-half

being collected, shaken with a little lime, soda, or potash, to neutralize any acetic acid, and decompose ethereal salts, and again distilled till one-half has passed over; the product is dilute spirit of wine. It may be still further concentrated or *rectified* by repeating this process of *fractional distillation*.

Both cane-sugar and grape-sugar yield alcohol by fermentation, the cane-sugar probably always passing into grape-sugar before the production of alcohol commences.

$$\underset{\text{Grape-sugar.}}{C_6H_{12}O_6} = \underset{\text{Alcohol.}}{2C_2H_5HO} + \underset{\text{Carbonic acid gas.}}{2CO_2}$$

Traces of several other substances are simultaneously produced (*vide* "*Fousel Oil*" in Index). By this reaction are formed the spirit of the various kinds of wine, beer, and liqueurs, such as Orange Wine (*Vinum Aurantii*, B. P.), made "by the fermentation of a saccharine solution, to which the fresh peel of the bitter orange has been added;" Sherry Wine (*Vinum Xericum*, B. P. and U. S. P.), the fermented juice of the grape; Port Wine (*Vinum Portense*, U. S. P.), Whiskey (*Spiritus Frumenti*, U. S. P.), containing from 48 to 56 per cent. of pure alcohol; *Spirit of Myrcia* or *Bay Rum* (*Spiritus Myrciæ*, U. S. P.), prepared by distilling rum with leaves of *Myrcia acris;* and others.

*Alcoholic drinks* vary much in strength. Cider or apple wine, cherry or pear wine, and good beer (ale and porter or stout) contain 4 to 6 per cent. of real alcohol; good light wines, both "red" and "white," 10 to 12 per cent.; good sherry and port, which are commonly "fortified," that is, contain added spirit, 16 or 18 per cent.; while "spirits" (gin, rum, brandy, whiskey, etc.) and "liqueurs" (ratafia, almond-flavored; maraschino, cherry-flavored; curaçoa, orange-flavored; chartreuse, a composite-flavored liqueur, etc.) are "under proof" or "over proof," terms explained in the next paragraph. The well-known effects of these fluids on the animal system would appear to be due primarily to alcohol, and, secondarily, to ethereal derivatives of alcohols. Some of the effect is also due to non-volatile substances, for beer from which all alcohol has been removed by ebullition still has a powerful influence on the human economy.

*Varieties of Alcohol.*—The weak spirit concentrated by distillation till it contains 84 per cent. by weight of pure alcohol is an ordinary article of commerce; its specific gravity at 60° is 0.8382. This is common *Spirit of Wine*, the *Spiritus Rectificatus* of the British Pharmacopœia. *Alcohol*, U. S. P., has sp. gr. 0.835: *Alcohol Fortius*, U. S. P., sp. gr. 0.817. The British official *Proof Spirit** (*Spiritus Tenuior*, B. P.) contains 49 per cent. by weight of

* *Proof spirit* is so termed from the fact that in olden times a proof of its strength was supposed to be afforded by moistening a small quantity of gunpowder and setting light to the spirit: if it fired the powder it was said to be "over proof;" if not, "under proof." The weakest spirit that would stand this test was what we should now describe as of sp. gr. 0.920.

alcohol, and is made by diluting 100 volumes of Rectified Spirit with water until the well-stirred product measures 156 volumes. Sixty volumes of water will be required for this purpose, the liquids occupying less bulk after than before admixture. In the language of the Excise authorities the rectified spirit of the Pharmacopœia would be described as "56 per cent. over proof" (56 per cent. O. P.); that is, 100 volumes contain as much alcohol as is present in 156 volumes of proof spirit. Obviously, proof spirit may be made by diluting with water rectified spirit of any other strength than that mentioned above. Thus 100 fluidounces of a spirit of "seventy over proof" may be diluted to 170, or the same quantity of a spirit of "fifty over proof" may be diluted to 150, and so on. The specific gravity of proof spirit at 60° is 0.920. (*Alcohol Dilutum*, U. S. P., has sp. gr. 0.941.)

*Empirical Formulæ.*—*Composition of Alcohol.*—Alcohol, by quantitative analysis, is found to contain the elements carbon, hydrogen, and oxygen in the following proportions:—

*Composition of Alcohol.*

| | | |
|---|---|---|
| Carbon . . . | 52.174 or ÷ 12 = | 4.348 or 2. |
| Hydrogen . . . | 13.043 or ÷ 1 = | 13.043 or 6. |
| Oxygen . . . | 34.783 or ÷ 16 = | 2.175 or 1. |
| | 100.000 | |

From centesimal numbers a formula is obtained in the usual way. Thus, on dividing these figures by the atomic weights of the respective elements (C = 12, H = 1, O = 16), and reducing the products to the simplest whole numbers, alcohol will be found to contain two atoms of carbon to every six of hydrogen and to every one of oxygen, and its possible or *empirical formula* to be $C_2H_6O$.

*Constitution of Alcohol.*—There is good reason to believe that alcohol is the hydrate of a basylous radical *ethyl* ($C_2H_5$ or Et); hence we derive the *rational formula* $C_2H_5HO$ or EtHO.

*Rational Formulæ.*—Rational formulæ are deduced by (1) ascertaining how much of the substance will combine with, displace, or play the part of, the atomic weight of a well-known element or radical. When this method cannot be applied, or in confirmation of it, processes of (2) reduction, (3) oxidation, (4) substitution, etc., are employed.

*Salts of Ethyl.*—Alcohol is, then, a body analogous in constitution to hydrate of potassium (KHO); and there are other compounds of ethyl analogous in constitution to ordinary inorganic salts, such as those of potassium. The oxide of ethyl ($Et_2O$) is common ether; the nitrite of ethyl ($EtNO_2$) is the chief body which, dissolved in spirit of wine, constitutes "sweet spirit of nitre;" the acid sulphate of ethyl ($EtHSO_4$), or sulphethylic or sulphovinic acid, is a liquid met with in the preparation of ether. The iodide (EtI), hydride (EtH), acetate ($Et\bar{A}$) and other salts are of considerable chemical interest, but not used in medicine.

*Absolute or Real Alcohol* ($C_2H_5HO$) may be prepared from spirit of wine by removing the water which the latter contains. This is accomplished, partially, by the agency of carbonate of potassium, and finally and entirely by recently burned quicklime. In operating on, say, one pint, 1½ ounce of dried carbonate of potassium is placed in a bottle that can be well closed, and frequently shaken during two days with the spirit. Meanwhile, about half a pound of good quicklime, if not already at hand, is made from 10 or 11 ounces of slaked lime by heating to redness in a covered crucible for half an hour. The spirit having been decanted from the denser aqueous solution of carbonate of potassium and placed in a quart flask, retort, or tin can, the lime, as soon as cold, is added, and the whole occasionally shaken during a day. The vessel is now placed in a saucepan or other bath containing water, quickly connected with a condenser (in the case of the flask or can by a bent tube and cork previously prepared; for absolute alcohol must not be exposed to air, or water in the form of moisture will be rapidly reabsorbed) and heat applied to the bath. Rejecting the first ounce or ounce and a half, as likely to contain traces of moisture absorbed from the air or apparatus, continue distillation until nothing more passes over, the water in the bath being kept just below the boiling-point (about 200° F.). These details are those of the British Pharmacopœia. Specific gravity 0.7938. Boiling-point 173° F.

*Tests.*—There are no specific tests for alcohol when mixed with complex matters. It is, however, easily isolated and concentrated by fractional distillation, and is then recognizable by conjoint physical and chemical characters. Thus its odor and taste are characteristic; it is lighter than water, volatile, colorless, and, when tolerably strong, inflammable, burning with an almost non-luminous flame; it readily yields aldehyd (see below) and acetic ether (*vide* p. 431), each of which has a characteristic odor; lastly, in presence of hot acid, alcohol reduces red chromate of potassium to a green salt of chromium.

According to Lieben 1 of alcohol in 2000 of water can be detected by adding to some of the warmed liquid a little iodine, a few drops of solution of soda, again warming gently, and setting aside for a time; a yellowish crystalline deposit of *iodoform* ($CHI_3$) is obtained. Under the microscope the latter presents the appearance of hexagonal plates or six-rayed and other varieties of stellate crystals.

$$C_2H_6O + 4I_2 + 6NaHO = CHI_3 + NaCHO_2 + 5NaI + 5H_2O.$$

Other alcohols, aldehyds, gum, turpentine, sugar, and several other substances give a similar reaction.

*Tests of purity.*—Oil or resin is precipitated on diluting spirit of wine with distilled water, giving an opalescent appearance to the mixture. The specific gravity should be 0.838. Fusel oil, aldehyd, and aldehydic acid are detected by nitrate of silver (*vide* Index, "Alcohol, test for purity of"). Water in absolute alcohol may be detected by adding to a small quantity a little highly dried sulphate of copper, which becomes blue ($CuSO_4,5H_2O$) if water is present,

but retains its yellowish-white anhydrous character ($CuSO_4$) if water be absent.

ALDEHYD ($C_2H_4O$).—Place together, in a capacious test-tube, or a flask, spirit of wine, black oxide of manganese, sulphuric acid, and water, and gently warm the mixture; aldehyd (*alcohol dehydrogenatus*), a highly volatile liquid, is immediately formed, and its vapor evolved, recognized by its peculiar, somewhat fragrant odor. Adapt a cork and rather long bent tube to the test-tube, and let some of the aldehyd slowly distil over into another test-tube, the condensing-tube being kept as cool as possible. Set the distillate aside for a day or two; the aldehyd will have nearly all disappeared, and acetic acid be found in the tube. Test the exposed liquid by litmus paper; it will be found to have an acid reaction: make it slightly alkaline by a drop or two of solution of carbonate of sodium, then boil to remove any alcohol and aldehyd present, add sulphuric acid, and notice the characteristic odor of the acetic acid evolved.

These experiments will enable the process of acetification described in connection with acetic acid to be more fully understood. Pure diluted alcohol is not oxidized by exposure to air; but in presence of fermentive matter, or vegetable matter undergoing decay or change, it is oxidized first to aldehyd and then to acetic acid.

In the above process the black oxide of manganese and sulphuric acid furnish nascent oxygen:—

$$\underset{\text{Black oxide of manganese.}}{MnO_2} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Sulphate of manganese.}}{MnSO_4} + \underset{\text{Oxygen. (atom).}}{O} + \underset{\text{Water.}}{H_2O}$$

The nascent oxygen then acts on the alcohol, just as the oxygen of the air acts on the alcohol in fermented infusion of malt, beer, or wine, giving aldehyd:—

$$\underset{\text{Alcohol.}}{C_2H_6O} + \underset{\text{Oxygen. (atom).}}{O} = \underset{\text{Aldehyd.}}{C_2H_4O} + \underset{\text{Water.}}{H_2O}$$

The aldehyd rapidly, even when pure (more rapidly when impure), absorbs oxygen and yields acetic acid:—

$$\underset{\text{Aldehyd.}}{2C_2H_4O} + \underset{\text{Oxygen.}}{O_2} = \underset{\text{Acetic acid.}}{2C_2H_4O_2}$$

*Tests.*—Aldehyd heated with solution of potash gives a brownish-yellow resinous mass of peculiar odor. Its aqueous solution reduces salts of silver, giving a mirror-like coating to the sides of a test-tube.

*Spirit of French Wine* (*Spiritus Vini Gallici*, B. P. and U. S. P.) or *Brandy* is a colored and flavored variety of alcohol distilled from French wine. Its color is that of light sherry, and is derived from the cask in which it has been kept, but it is commonly deepened by

the addition of burnt sugar. Its taste is due to the volatile flavoring constituent of the wine, often increased by the addition of artificial essences. It should contain from 48 to 56 per cent. of alcohol.

## QUESTIONS AND EXERCISES.

753. Write a few sentences on the formation, purification, and concentration of alcohol, and explain the difference between Rectified Spirit, Proof Spirit, and Absolute Alcohol.

754. What quantity of water must be added to one gallon of spirit of wine, 56 degrees over proof, to convert it into proof spirit?

755. To what volume must 5 pints of spirits of wine of 53 degrees be diluted before it becomes proof spirit?—*Ans.* 7 pints, 13 ounces.

756. State the specific gravity of proof spirit.

757. Show how the formula of alcohol is obtained from its centesimal composition:—

| | |
|---|---|
| Carbon | 52.174 |
| Hydrogen | 13.043 |
| Oxygen | 34.783 |
| | 100.000 |

758. Give the formulæ of some of the salts of ethyl.

759. By what processes may pure hydrate of ethyl be obtained?

760. Enumerate the characters of alcohol.

761. Mention a chemical test to distinguish rectified spirit from absolute alcohol.

762. From the formula of aldehyd calculate back its composition in 100 parts.

763. What is the relation of aldehyd to alcohol and to acetic acid?

764. Whence is brandy obtained, and to what are due its color and flavor?

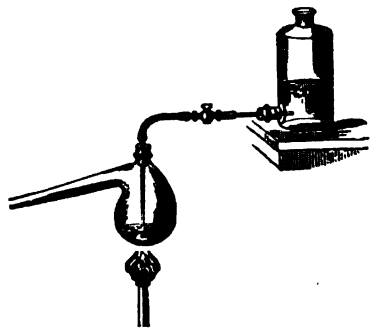
## ETHER, OR OXIDE OF ETHYL.

Formula $C_4H_{10}O$, or $(C_2H_5)_2O$, or $Et_2O$.

*Experimental Process.*— Into a capacious test-tube put a small quantity of spirit of wine and about half its bulk of sulphuric acid, mix and gently warm; the vapor of ether, recognized by its odor, is evolved. Adapt a cork and long bent tube to the test-tube, and slowly distil over the ether into another test-tube. Half the original quantity of alcohol now placed in the generating-tube will again give ether; and this operation may be repeated many times.

*On the large scale*, and according to the following official process (*Æther*, B. P. and U. S. P.), the addition of alcohol, instead of being intermitting, is continuous, a tube conveying alcohol from a reservoir into the generating-vessel. Mix 10 fluidounces of sulphuric acid with 12 fluidounces of rectified spirit in a glass retort or flask capable of containing at least two pints, and, not allowing the mixture to

Fig. 49.



cool, connect the retort or flask by means of a bent glass tube with a Liebig's condenser, and distil with a heat sufficient to maintain the liquid in brisk ebullition. If a thermometer also be inserted in the tubulars of the retort, or through the cork of the flask, the temperature may be still more carefully regulated—between 284° and 290° F. As soon as the ethereal fluid begins to pass over, supply fresh spirit in a continuous stream, and in such quantity as to about equal the volume of the fluid which distils. For this purpose use a tube furnished with a stopcock to regulate the supply, connecting one end of the tube with a vessel containing the spirit supported above the level of the retort or flask, and passing the other end through the cork of the flask into the liquid. When a total of 50 fluidounces of spirit has been added, and 42 fluidounces of ether have distilled over, the process may be stopped.

To *partially purify* the liquid, dissolve 10 ounces of chloride of calcium in 13 ounces of water, add half an ounce of lime, and agitate the mixture in a bottle with the impure ether. Leave the mixture at rest for ten minutes, pour off the light supernatant fluid, and distil it with a gentle heat until a glass bead of specific gravity 0.735 placed in the receiver begins to float. The ether and spirit retained by the chloride of calcium and by the residue of each rectification may be recovered by distillation and used in a subsequent operation.

*Explanation of Process.*—On the addition of sulphuric acid to alcohol in equal volumes, one molecule of each react and give a molecule of sulphethylic acid and one of water:—

$$\underset{\text{Alcohol.}}{EtHO} + \underset{\substack{\text{Sulphuric}\\\text{acid.}}}{H_2SO_4} = \underset{\substack{\text{Sulphethylic}\\\text{acid.}}}{EtHSO_4} + \underset{\text{Water.}}{H_2O}$$

More alcohol then gives ether and sulphuric acid by the reaction of one molecule of the alcohol on one of sulphethylic (ethylsulphuric) acid:—

$$\underset{\text{Alcohol.}}{EtHO} + \underset{\substack{\text{Sulphethylic}\\ \text{acid.}}}{EtHSO_4} = \underset{\text{Ether.}}{Et_2O} + \underset{\substack{\text{Sulphuric}\\ \text{acid.}}}{H_2SO_4}$$

The water of the first reaction and the ether of the second distil over, while the sulphuric acid, as fast as liberated, is attacked by alcohol and reconverted into sulphethylic acid:—

$$\underset{\text{Alcohol.}}{EtHO} + \underset{\substack{\text{Sulphuric}\\ \text{acid.}}}{H_2SO_4} = \underset{\substack{\text{Sulphethylic}\\ \text{acid.}}}{EtHSO_4} + \underset{\text{Water.}}{H_2O}$$

so that the sulphuric acid originally employed finally remains in the retort in the form of sulphethylic acid. The effect, however, of a small quantity of sulphuric acid in thus converting a large quantity of alcohol into ether is limited, secondary reactions occurring to some extent after a time.

*Properties.*—Pure ether is gaseous at temperatures above 95° F.; hence the condensing-tubes employed in its distillation must be kept as cool as possible. At all ordinary temperatures it rapidly evaporates, absorbing much heat from the surface on which it is placed. A few drops evaporated consecutively from the back of the hand produce great cold; if blown in the form of spray, the cooling effect is so rapid and intense as to produce local anæsthesia. Its vapor is very heavy, more than twice and one-half that of air and nearly forty times that of hydrogen ($H_2 = 2$; $C_4H_{10}O = 74$; or as 1 to 37). In a still atmosphere, therefore, it will flow a considerable distance along a table or floor before complete diffusion occurs; the vapor is also highly inflammable; hence the importance of keeping candle and other flames at a distance during manipulations with ether.

*Purification.*—To imitate the process of partial purification above described, add to the small quantity of ether obtained in the foregoing operation a strong solution of chloride of calcium and a little slaked lime; the latter absorbs any sulphurous acid that may have been produced by secondary decompositions, while the former absorbs water; on shaking the mixture and then setting aside for a minute or two, the ether will be found floating on the surface of the solution of chloride of calcium.

This ether, redistilled until the distillate has a sp. gr. not higher than 0.735 (0.750, U. S. P.) and boiling-point not higher than 105° F., is the ether of the British Pharmacopœia. It still contains about 8 per cent. of alcohol. The latter may be removed by well shaking the ether with half of its bulk of water, setting aside, separating the floating ether and again shaking it with water; alcohol is thus washed out. This washed ether containing water (for water and ether are to some extent soluble the one in the other; 50 measures agitated with an equal volume of water are reduced to 45 by an absorption of 10 per cent.) is next placed in a retort with solid chlo-

ride of calcium and a little caustic lime, and once more distilled; pure dry ether (*Æther Purus*, B. P., *Æther Fortior*, U. S. P.) results. Sp. gr. not exceeding 0 720 (0.728, U. S. P.).

*Spiritus Ætheris*, B. P., is a mixture of common ether (*Æther*, B. P.) with twice its bulk of rectified spirit.

## NITROUS ETHER, OR NITRITE OF ETHYL.

Formula $C_2H_5NO_2$ or $EtNO_2$.

*Process.*—To a third of a test-tubeful of rectified spirit add about a tenth of its bulk of sulphuric acid, rather more of nitric acid, and some copper-wire or turnings, and warm the mixture as soon as ebullition commences; the vapor of nitrous ether is evolved, recognized by its odor. A long bent tube, kept cool, may be adapted by a perforated cork to the test-tube, and thus a few drops of impure nitrous ether be condensed and collected.

*The above process conducted on a larger scale*, with definite quantities of materials, temperature regulated by a thermometer, and a well-cooled condenser, etc. (see p. 126), is the official (Redwood's) process for the preparation of a concentrated solution of nitrous ether in spirit; diluted with nearly three times its bulk of rectified spirit it forms the "sweet spirit of nitre" (*Spiritus Ætheris Nitrosi*, B. P. and U S. P.) of pharmacy.

"Take of

| | |
|---|---|
| Nitric Acid . . . . . | 3 fluidounces, |
| Sulphuric Acid . . . . . | 2 fluidounces, |
| Copper, in fine wire (about No. 25) | 2 ounces, |
| Rectified Spirit . . . . . | a sufficiency. |

"To one pint of the spirit add gradually the sulphuric acid, stirring them together; then add, in the same way, two and a half ounces of the nitric acid. Put the mixture into a retort or other suitable apparatus, into which the copper has been introduced, and to which a thermometer is fitted. Attach now an efficient condenser, and, applying a gentle heat, let the spirit distil at a temperature commencing at 170° and rising to 175°, but not exceeding 180°, until 12 fluidounces have passed over and been collected in a bottle kept cool, if necessary, with ice-cold water; then withdraw the heat, and, having allowed the contents of the retort to cool, introduce the remaining half ounce of nitric acid, and resume the distillation as before, until the distilled product has been increased to 15 fluidounces. Mix this with two pints of the rectified spirit, or as much as will make the product correspond to the tests of specific gravity and percentage of liquid separated by chloride of calcium (*vide infra*). Preserve it in well-closed vessels."

Disregarding secondary products (including much aldehyd), the following equation probably represents decompositions that occur in

the operation. The main point in the reaction is the reduction of the nitric to the nitrous radical by the indirect agency of the copper.

$$\underset{\text{Alcohol.}}{EtHO} + \underset{\text{Nitric acid.}}{HNO_3} + \underset{\text{Sulphuric acid.}}{H_2SO_4} + \underset{\text{Copper.}}{Cu} = \underset{\text{Nitrous ether.}}{EtNO_2} + \underset{\text{Water.}}{2H_2O} + \underset{\text{Sulphate of copper.}}{CuSO_4}$$

*Properties.*—Spirit of Nitrous Ether "is transparent and nearly colorless, with a very slight tinge of yellow, mobile, inflammable, of a peculiar penetrating apple-like odor, and sweetish, cooling, sharp taste. Specific gravity, 0.845. It effervesces, feebly, or not at all, when shaken with a little bicarbonate of soda" (showing absence of appreciable quantities of free nitrous, acetic, or other acids). The aldehyd in it may be detected by the potash test (see page 426). The great tendency of aldehyd to become converted into acetic acid by the absorption of oxygen from the air renders Spirit of Nitrous Ether unstable, and pharmacists are obliged to neutralize such acid, generally by bicarbonate of potassium, before adding it to medicines in which are ingredients incompatible with acids.

*Test.*—The nitrous radical may be detected by adding sulphate of iron and sulphuric acid to some of the spirit of nitrous ether, a brown or black compound being produced, already explained in connection wth nitric acid (p. 289).

*Test of Strength.*—To some of the official Spirit of Nitrous Ether add twice its bulk of a saturated solution of chloride of calcium and mix the liquids; on setting aside, an ethereal fluid will rise to the surface. If the spirit of nitrous ether be of official strength, not less than 2 per cent. of its volume will thus separate, indicating the presence of 10 per cent. of the ethereal fluid, 8 per cent. being said to be still in solution. A graduated tube is obviously most convenient for this purpose.

*Spiritus Ætheris Nitrosi*, U. S. P., has the specific gravity 0.837, and contains five per cent. of nitrous ether.

## Acetic Ether, or Acetate of Ethyl.

To a little dried acetate of sodium, in a test-tube, add a small quantity of rectified spirit of wine and some sulphuric acid, and, adapting a long bent tube in the usual manner, heat the test-tube and so distil over acetic ether—which may be collected in another test-tube kept cool by partial immersion in cold water.

The official proportions (*Æther Aceticus*, B. P.) are 8 parts of dried acetate of sodium, 5 of spirit, and 10 of acid. It is purified from any water by shaking in a bottle with fused chloride of calcium, and, after twenty-four hours, rectifying. "Sp. gr. 0.910. Boiling-point 166° F."

$$\underset{\text{Hydrate of ethyl.}}{C_2H_5HO} + \underset{\text{Acetate of sodium.}}{NaC_2H_3O_2} + \underset{\text{Sulphate of hydrogen.}}{H_2SO_4} = \underset{\text{Acetate of ethyl.}}{C_2H_5C_2H_3O_2} + \underset{\text{Sulphate of sodium and hydrogen.}}{NaHSO_4} + \underset{\text{Water.}}{H_2O}$$

IODIDE OF ETHYL (EtI) may be prepared by mixing amorphous phosphorus with *absolute* alcohol and adding iodine gradually.

$$\underset{\text{Alcohol.}}{5EtHO} + \underset{\substack{\text{Iodide of}\\\text{phosphorus.}}}{PI_5} = \underset{\substack{\text{Iodide of}\\\text{ethyl.}}}{5EtI} + \underset{\substack{\text{Phosphoric}\\\text{acid.}}}{H_3PO_4} + \underset{\text{Water.}}{H_2O}$$

The reaction at first proceeds rapidly, and is complete after the mixture has been set aside for a few hours. The iodide of ethyl may then be isolated by careful distillation, freed from any excess of iodine by washing with a very small quantity of solution of potash or soda, washed with water, dried over chloride of calcium, and again distilled. It should be kept in a dark place, as light favors decomposition and liberation of iodine.

*Bromide of ethyl* ($C_2H_5Br$), or *hydrobromic ether*, is similarly prepared.

ETHYL.—This gaseous radical, $(C_2H_5)_2$ or $Et_2$, is obtained on digesting together, at about 250° F., in a strong sealed tube, dry freshly granulated zinc with iodide of ethyl (Frankland).

$$\underset{\text{Zinc.}}{Zn} + \underset{\substack{\text{Iodide of}\\\text{ethyl.}}}{2EtI} = \underset{\substack{\text{Iodide of}\\\text{zinc.}}}{ZnI_2} + \underset{\text{Ethyl.}}{Et_2}$$

On cautiously opening the tube the ethyl escapes, and may be ignited or collected over water. There remains with the iodide of zinc a body termed by Frankland *zinc-ethyl* ($ZnEt_2$); it is a spontaneously inflammable liquid, but may easily be distilled and otherwise manipulated if a few simple precautions be observed. If water be allowed to flow down the tube, the solid compound of iodide of zinc and zinc-ethyl will be decomposed, a gas, *hydride of ethyl* (EtH), resulting, which also may be inflamed or collected over water:—

$$ZnEt_2 + 2H_2O = Zn2HO + 2EtH.$$

---

## QUESTIONS AND EXERCISES.

765. Describe the official process for the preparation of Ether, giving equations.

766. Offer a physical explanation of the mode of producing local anæsthesia.

767. How is commercial ether purified?

768. Explain Redwood's process for the preparation of "sweet spirit of nitre."

769. Give the properties of spirit of nitrous ether.

770. By what method is the strength of "sweet spirit of nitre" estimated?

771. How is iodide of ethyl made?

772. Adduce evidence of the existence of ethyl.

## OTHER ALCOHOL RADICALS AND THEIR SALTS.

What has been stated concerning the chemistry of ethyl and its compounds may be applied to other radicals known to exist, some of the compounds of each of which are of common occurrence. These basylous radicals are closely related to each other, to hydrogen, and to the metals. Their formulæ in the combined state may be built up by successive additions of $CH_2$, thus:—

| | |
|---|---|
| Hydrogen . . . . . | H |
| Methyl . . . . . | $CH_3$, or Me |
| Ethyl . . . . . | $C_2H_5$, or Et |
| Propyl (or Trityl) . . . . . | $C_3H_7$, or Pr |
| Butyl (or Tetryl) . . . . . | $C_4H_9$, or Bu |
| Amyl . . . . . | $C_5H_{11}$, or Ay |
| Caproyl (or Hexyl) . . . . . | $C_6H_{13}$, or Cp |

The above list is an illustration of an *homologous* series (from ὁμός, *homos*, the same, and λόγος, *logos*, proportion) of compounds. It will be observed that the relation of the number of hydrogen atoms to carbon is twice as many with one added; hence the series is often termed the $C_nH_{2n+1}$ series ($n$ = any number). The oxides of these radicals are known as *ethers*, their hydrates *alcohols*, their compounds with the acetic and similar acidulous radicals *ethereal salts*. Every alcohol furnishes a body corresponding to the aldehyd of spirit of wine, the class being termed *aldehyds;* each also yields an acid corresponding with acetic acid. Any one of these classes constitutes an homologous series. Or, taking the hydride, oxide, hydrate, acid, of any single radical, we get a *heterologous* (ἕτερος, *heteros*, another) series of compounds. Hydride of methyl (MeH or $CH_3H$) is ordinary *marsh-gas*, *fire-damp*, or *light carburetted hydrogen;* it is a diluent or non-luminiferous constituent of ordinary coal-gas to the extent of 30 or 40 per cent.;* *formic acid*, the acid of the methyl

* *Coal-gas.*—The other diluents, or vehicles for the conveyance of illuminating vapors of coal-gas, are hydrogen (40 to 50 per cent.) and carbonic oxide (6 to 7 per cent.). The illuminating constituents are olefiant gas (p. 447) and its homologues existing to the extent of from 5 to 7 per cent. Hydrocarbons normally fluid but kept in the vaporous condition by the diluents also contribute materially to the illuminating power of gas. The impurities are nitrogen air carbonic acid, bisulphide of carbon $CS_2$ (a volatile liquid easily made from its elements and rendered almost scentless by digesting with copper turnings), and other badly smelling sulphur compounds. Upwards of fifty distinct chemical substances have been obtained from the solid, liquid, and gaseous products of the destructive distillation of coal.

series; *butyric acid* the acid of the butyl series; *sulphocyanate of butyl*, the essential oil of horseradish; *valerianic acid*, the acid of the amyl series.

*Homologous and Heterologous Series of the* C $H_{2n+1}$ *Radicals.*

| Radicals (free). | Hydrides. | Oxides (or ethers). | Hydrates (or alcohols). | Aldehyds. | Acids. |
|---|---|---|---|---|---|
| $(CH_3)_2$ | $CH_3H$ | $(CH_3)_2O$ | $CH_3HO$ | $CH_2O$? | $CH_2O_2$ |
| $(C_2H_5)_2$ | $C_2H_5H$ | $(C_2H_5)_2O$ | $C_2H_5HO$ | $C_2H_4O$ | $C_2H_4O_2$ |
| $(C_3H_7)_2$ | $C_3H_7H$ | $(C_3H_7)_2O$ | $C_3H_7HO$ | $C_3H_6O$ | $C_3H_6O_2$ |
| $(C_4H_9)_2$ | $C_4H_9H$ | $(C_4H_9)_2O$ | $C_4H_9HO$ | $C_4H_8O$ | $C_4H_8O_2$ |
| $(C_5H_{11})_2$ | $C_5H_{11}H$ | $(C_5H_{11})_2O$ | $C_5H_{11}HO$ | $C_5H_{10}O$ | $C_5H_{10}O_2$ |
| etc. | etc. | etc. | etc. | etc. | etc. |

## QUESTIONS AND EXERCISES.

773. Mention several radicals homologous with ethyl, and give their formulæ.

774. Define ethers, hydrides, alcohols, ethereal salts, aldehyds.

775. What is the difference between homologous and heterologous series?

776. Give the systematic name of *fire-damp*.

777. Enumerate the chief constituents of coal-gas.

778. State the formulæ of formic, butyric, and valerianic acids.

779. Write the formulæ of butyl, its hydride, ether, alcohol, aldehyd, and acid.

## METHYLIC ALCOHOL.

Methylic Alcohol ($CH_3HO$, or MeHO), Wood-Spirit, or Pyroxylic Spirit, or Wood-Naphtha, is a product of the destructive distillation of wood. Spirit of wine containing 10 per cent. of wood-spirit constitutes ordinary *methylated spirit*, a spirit issued duty free, for the use of manufacturers, the methylic alcohol not interfering with technical applications. From its nauseous taste and odor, however, it cannot take the place of gin, brandy, or other spirit; hence, while industry is benefited, intemperance is discouraged and the revenue not injured.

*Detection of Methylic Alcohol in presence of Ethylic Alcohol.*—Three or four methods have been proposed for the detection

of methylated spirit in various liquids; that open to least objection is by J. T. Miller. For the application of the test to tinctures and similar spirituous mixtures, some of the spirit is first separated by distilling off a drachm or so from about half an ounce of the liquid placed in a small flask or test-tube having a long bent tube attached. Into a similar apparatus put 30 grains of powdered red chromate of potassium, half an ounce of water, 25 minims of strong sulphuric acid, and 30 or 40 minims of the spirit to be tested. Set the mixture aside for a quarter of an hour, and then distil nearly half a fluidounce. Place the distillate in a small dish, add a very slight excess of carbonate of sodium, boil down to about a quarter of an ounce, add enough acetic acid to impart a distinct but feeble acid reaction, pour the liquid into a test-tube, add a grain of nitrate of silver dissolved in about 30 drops of water, and heat gently for a couple of minutes. If the liquid then merely darkens a little, but continues quite translucent, the spirit is free from methylic alcohol; but if a copious precipitate of dark-brown or black metallic silver separates, and the tube, after being rinsed out and filled with clear water, has a distinct film of silver, which appears brown by transmitted light (best seen by holding it against white paper), the spirit is methylated.

*Explanation.*—This test depends for its action on the reducing-powers of formic acid. In the above operation the ethylic alcohol becomes oxidized to acetic acid (the natural acid of the ethyl series), which does not reduce silver salts, a minute quantity only of formic acid being produced, while the methylic alcohol yields formic acid (the natural acid of the methyl series) in a comparatively large quantity. Aldehyd, which is also a reducing agent, is simultaneously produced, but removed in the subsequent ebullition with carbonate of sodium.

*Methylated Sweet Spirit of Nitre.*—The preparation of spirit of nitrous ether from methylated spirit is illegal in Great Britain, but, nevertheless, is occasionlly practised. For the detection of methylic alcohol in this liquid, Mr. Miller suggests the following modification of the above process.

Shake about an ounce of the sample with 20 or 30 grains of anhydrous carbonate of potassium, and, if needful, add fresh portions of the salt until it ceases to be dissolved, then pour off the supernatant spirit. This serves to neutralize acid and to remove water, in which some samples are remarkably rich. Introduce half a fluidounce of the spirit into a small flask; add 150 grains of anhydrous chloride of calcium in powder, and stir well together; then, having connected the flask with a condenser, place it in a bath of boiling water, and distil a

fluidrachm and a half, or continue the distillation until scarcely anything more comes over. The operation is rather slow, but needs little attention, and should be done thoroughly. The distillate contains nearly the whole of the nitrous ether and other interfering substances, while in the retort there remains a non-volatile compound of chloride of calcium and methylic alcohol, if the latter be present. Now add to the contents of the flask a fluidrachm of water, which decomposes the compound just referred to, and draw over the half drachm of spirit required for testing. Add to it the usual oxidizing solution composed of 30 grains of red chromate of potassium, 25 minims of strong sulphuric acid, and half an ounce of water; let the mixture stand a quarter of an hour, then distil half a fluidounce. Treat the distillate with a slight excess of carbonate of sodium, boil rapidly down to two fluidrachms, and drop in, cautiously, enough acetic acid to impart a faint acid reaction; pour the liquid into a test-tube about three-quarters of an inch in diameter; add two drops of *diluted* acetic acid, and one grain of nitrate of silver in half a drachm of pure water; apply heat, and boil gently for two minutes. If the spirit is free from methylic alcohol, the solution darkens and often assumes transiently a purplish tinge, but continues quite translucent, and the test-tube, after being rinsed out and filled with water, appears clean or nearly so. But if the spirit contains only 1 per cent. of methylic alcohol, the liquid turns first brown, then almost black and opaque, and a film of silver, which is brown by transmitted light, is deposited on the tube. When the sample is methylated to the extent of 3 or 4 per cent., the film is sufficiently thick to form a brilliant mirror. To insure accuracy, the experiments should be performed by daylight.

## CHLOROFORM.

Formula $CHCl_3$.

*Process.*—Should the necessary appliances be at hand, a small quantity of this liquid may easily be prepared by the official process. One fluidounce and a half of spirit and 24 of water are placed in a retort or flask of at least a quart capacity; 8 oz. of chlorinated lime and 4 of slaked lime are added, the vessel connected with a condenser, and the mixture heated until distillation commences, the source of heat then being withdrawn. The condensed liquid should fall into a small flask containing water, at the bottom of which about a drachm of chloroform will slowly collect.

*Explanation of Process.*—The hypochlorite of calcium (Ca2ClO) believed to be present in the chlorinated lime (see remarks in connection with the latter, p. 113) readily yields up oxygen and chlorine to organic substances, the calcium being liberated as hydrate, $4(Ca2ClO) + 4H_2O = 4(Ca2HO) + 2O_2 + 4Cl_2$. The alcohol used in making chloroform is thus reduced first to aldehyd:—

$$\underset{\text{Alcohol.}}{2C_2H_6O} + \underset{\text{Oxygen.}}{O_2} = \underset{\text{Aldehyd.}}{2C_2H_4O} + \underset{\text{water.}}{2H_2O}$$

The action of chlorine on aldehyd then probably gives chloral (*chlor-al*dehyd):—

$$\underset{\text{Aldehyd.}}{C_2H_4O} + \underset{\text{Chlorine.}}{3Cl_2} = \underset{\text{Chloral.}}{C_2HCl_3O} + \underset{\text{Hydrochloric acid.}}{3HCl}$$

The hydrochloric acid being at once neutralized by some of the liberated hydrate of calcium to form chloride of calcium and water, more freed hydrate of calcium and chloral give formate of calcium and chloroform.

$$\underset{\text{Chloral.}}{2C_2HCl_3O} + \underset{\text{Hydrate of calcium.}}{Ca2HO} = \underset{\text{Formate of calcium.}}{Ca2CHO_2} + \underset{\text{Chloroform.}}{2CHCl_3}$$

Or, neglecting the probable steps in the process, and regarding only the materials and the products, 4 molecules of alcohol and 8 of hypochlorite of calcium give two of chloroform, 3 of formate of calcium, 5 of chloride of calcium, and 8 of water, thus:—

$$\underset{\text{Alcohol.}}{4C_2H_6O} + \underset{\text{Hypochlorite of calcium.}}{8CaCl_2O_2} = \underset{\text{Chloroform.}}{2CHCl_3} + \underset{\text{Formate of calcium.}}{3(Ca2CHO_2)} + \underset{\text{Chloride of calcium.}}{5CaCl_2} + \underset{\text{Water.}}{8H_2O}$$

The hydrate of calcium placed in the generating-vessels is not essential, but is useful in preventing secondary decompositions, the hydrate of calcium obtainable from the reaction being insufficient for this purpose.

*Constitution.*—Chloroform is sometimes considered to be the chloride of a trivalent radical methenyl (CH), the first member of a series $C_nH_{2n-1}$. Glycerine is the hydrate of another member—*glyceryl* or *propenyl*, $C_3H_5$ (p. 448).

Chloroform may also be regarded as the chloride of dichlor-methyl; it may be formed from methylic compounds, thus:—

$$\underset{\text{Methylic alcohol.}}{2CH_4O} + \underset{\text{Chlorinated lime.}}{2(CaCl_2,CaCl_2O_2)} = \underset{\text{Chloroform.}}{2CHCl_3} + \underset{\text{Oxychloride of calcium.}}{CaCl_2,3CaO} + \underset{\text{Water.}}{3H_2O}$$

Chlorine converts it into tetrachloride of carbon, completing a series of *substitution products* of chloride of methyl.

$$\underset{\text{Chloride of methyl.}}{C\left\{\begin{matrix}H\\H\\H\end{matrix}\right\}Cl} \quad \underset{\text{Chloride of mono-chlor-methyl.}}{C\left\{\begin{matrix}H\\H\\Cl\end{matrix}\right\}Cl} \quad \underset{\text{Chloride of di-chlor-methyl.}}{C\left\{\begin{matrix}H\\Cl\\Cl\end{matrix}\right\}Cl} \quad \underset{\text{Chloride of tri-chlor-methyl.}}{C\left\{\begin{matrix}Cl\\Cl\\Cl\end{matrix}\right\}Cl} \text{ or } \underset{\text{Tetrachloride of carbon.}}{CCl_4}$$

The chloride of mono-chlor-methyl, under the name of dichloride

of methylene, has been used as an anæsthetic. It may be obtained by the action of nascent hydrogen on chloroform.

*Chloroform is purified* by shaking it with pure sulphuric acid containing no trace of nitric acid, which chars and removes hydrocarbons, but does not affect chloroform. It is freed from any trace of acid by agitation with carbonate of sodium, and from moisture by distillation with lime.

*Properties.*—The sp. gr. of purified chloroform is 1.500. It is liable to slowly decompose when exposed to air and light. To render it stable a minute amount of alcohol is necessary; hence the specific gravity of medicinal chloroform is about 1.497 (1.480 U. S. P.). It readily and entirely volatilizes with characteristic odor at common temperatures. It has a sweetish taste, is limpid, colorless, soluble in alcohol and ether, and slightly in water, but burns with a sluggish green smoky flame.

*Aqua Chloroformi*, B. P., the official chloroform water, is made by shaking 1 fluidrachm of chloroform with 25 ounces of distilled water till dissolved.

*Iodoform* (*Iodoformum*, U. S. P.), analogous in constitution to chloroform, the iodine taking the place of the chlorine ($CHI_3$), is prepared by mixing in a retort 2 parts of carbonate of potassium, 2 parts of iodine, 1 part of alcohol, and 5 of water, heating till colorless, and then pouring into a beaker and allowing to settle. The Iodoform is deposited in yellow scales, which are collected on a filter, washed thoroughly with water, and dried between filtering paper.

Iodoform appears in the shape of yellow, shining six-sided scales with a saffron-like odor; is volatile at ordinary temperatures, and at a temperature above 250° is decomposed, giving off violet vapors. Almost insoluble in water, but more so in alcohol, ether, and the fixed and volatile oils.

## CHLORAL AND CHLORAL HYDRATE.

*Process.*—Pass a rapid stream of dry chlorine into pure absolute alcohol so long as absorption occurs. During the first hour or two the alcohol must be kept cool, afterwards gradually warmed till ultimately the boiling-point is reached. The preparation of a considerable quantity occupies several days. The crude product is mixed with three times its volume of oil of vitriol and distilled, again mixed with a similar quantity of oil of vitriol and again distilled, and finally rectified from quicklime.

The formation of chloral would at first sight seem to be due to the production from the alcohol ($C_2H_6O$) of aldehyd ($C_2H_4O$), through the removal of hydrogen by the chlorine, and the substitution of chlorine for hydrogen in the aldehyd ($C_2H_4O$) with formation of *chlor-aldehyd* or *chloral* ($C_2HCl_3O$). But the reactions are far more complicated, being somewhat as follows. Aldehyd and hydrochloric acid are first formed; these with some of the alcohol give monochlo-

rinated ether, $\left.\begin{matrix}C_2H_5\\C_2H_4Cl\end{matrix}\right\}O$, which with more chlorine yields tetrachlorinated ether, $\left.\begin{matrix}C_2H_5\\C_2HCl_4\end{matrix}\right\}O$, and this in presence of water furnishes chloral, some alcohol, and some hydrochloric acid (Wurtz and Vogt).

*Properties.*—The formula of chloral is $C_2HCl_3O$. It is a colorless liquid, of oily consistence. Sp. gr. 1.502. Boiling-point 201.2° F. Its vapor has a penetrating smell, and is somewhat irritating to the eyes. Mixed with water, heat is disengaged, and solid white, crystallizable, *hydrous chloral* (*Chloral*, U. S. P.), or what is more generally, though somewhat irregularly, termed *chloral hydrate* ($C_2HCl_3O,H_2O$) or *hydrate of chloral* (*Chloral Hydras*, B. P.), results. The latter fuses at 110.8° and boils at 203° F. It sublimes as a white crystalline powder. Both chloral and chloral hydrate are soluble in water, alcohol, ether, and oils. Oils and fats are, also, soluble in chloral hydrate. The aqueous solution should be neutral, and give no reaction with nitrate of silver. Chloral is said sometimes to undergo a spontaneous change into an opaque white isomeric modification, insoluble in water, alcohol, or ether, but convertible by prolonged contact with water or by distillation into the ordinary condition. By action of weak alkalies chloral first yields formate of the alkali-metal and chloroform:—

$$C_2HCl_3O + KHO = KCHO_2 + CHCl_3.$$

Chloral, or rather strong aqueous solution of chloral hydrate (3 in 4), injected beneath the skin yields nascent chloroform by action of the alkali of the blood, and produces narcotic effects (Liebreich, Personne). Chloroform itself admits of similar hypodermic use (Richardson). If administered by the stomach, thirty to eighty grains of solid hydrate are required. The final products of the reaction of the chloroform and blood are formate and chloride of sodium. A spirituous solution of potash effects the same transformation.

$$CHCl_3 + 4KHO = KCHO_2 + 3KCl + 2H_2O.$$

Solution of ammonia and moist hydrate of calcium, as well as weak solutions of fixed alkalies, convert hydrate of chloral into formate of the metal and chloroform. The reaction with the slaked lime being especially definite and complete (Wood), it may be employed in ascertaining the richness of a sample of commercial chloral hydrate in chemically pure chloral hydrate.

$$\underbrace{2(C_2HCl_3O,H_2O)}_{331} + Ca2HO = \underbrace{2CHCl_3}_{239} + Ca2CHO_2 + 2H_2O$$

From the foregoing equation and molecular weights it is obvious that 100 grains of hydrate of chloral, if kept dry, will yield by distillation with 30 grains of slaked lime and an ounce of distilled water (in a small flask and long bent tube kept cool by moistened paper) 72.2 grains of chloroform, by weight, or (the sp. gr. of chloroform being taken at 1.497) 47.56 grains by measure, or about 52 minims. The official hydrate of chloral "should yield not less than 70 grains of chloroform."

*Pure* Chloral Hydrate.—Liebreich, who first proposed the use of chloral hydrate, gives the following as the characteristics of a pure article: Colorless, transparent crystals. Does not decompose by the action of the atmosphere, does not leave oily spots when pressed between blotting-paper, affects neither cork nor paper. Smells agreeably aromatic, but a little pungent when heated. Taste bitter, astringent, slightly caustic. Seems to melt on rubbing between the fingers. Dissolves in water like candy without first forming oily drops; and the solution is neutral or faintly acid to test-paper. Dissolves in bisulphide of carbon, petroleum, ether, water, alcohol, oil of turpentine, etc. Its solution in chloroform gives no color when shaken with sulphuric acid. Boiling-point 203° to 205° F., and volatilizes without residue. Distilled with sulphuric acid, the chloral should pass over at 205° to 207° F. Melting-point 133° to 136° F.; again solidifying at about 120° F. Gives no chlorine reaction on treating the solution in water (acidulated by nitric acid) with nitrate of silver.

*Impure* Chloral Hydrate.—Yellowish, cloudy. Decomposes; leaves spots by pressing between blotting-paper; decomposes corks and paper of the packing. Smells pungent and irritating; on opening the bottles sticky and often emits fumes. Taste strongly caustic. With water forms oily drops or is partially insoluble. Boils at a higher temperature. On treating it with sulphuric acid turns brown, with formation of hydrochloric acid. Gives chlorine reaction on treating the solution in water (acidulated by nitric acid) with nitrate of silver.

*Alcoholates of chloral* are obtained on combining alcohols with chloral.

*Bromal* ($C_2HBr_3O$), *hydrate of bromal* ($C_2HBr_3O,H_2O$), and *alcoholates of bromal* are produced when bromine instead of chlorine attacks alcohol. *Iodal* ($C_2HI_3O$) also exists.

*Butyl chloral* ($C_4H_5Cl_3O$), originally, but erroneously termed

*croton chloral*, is a product of the action of chlorine on aldehyd. Its name expresses its constitution; it is *chlorinated butyric aldehyd;* ordinary chloral being chlorinated vinic aldehyd.

## AMYLIC ALCOHOL.

AMYLIC ALCOHOL (*Alcohol Amylicum*, B. P. and U. S. P.) ($C_5H_{11}HO$, or AyHO) is a constant accompaniment of ethylic or common alcohol ($C_2H_6HO$, or EtHO) when the latter is prepared from sugar which has been derived from starch; hence the name, from *amylum*, starch. The sugar of potato-starch yields a considerable quantity; hence the alcohol is often called *potato-oil*. It is also termed *fousel-oil* or *fusel-oil* (from φυω, *phuō*, to produce), in allusion to the circumstance that the supposed oil is not simply educed from a substance already containing it, as is usually the case with oils, but is actually produced during the operation. It was described as oil probably because it resembled oil in not readily mixing with water; but it is soluble to some extent in water, and is a true spirit, homologous with spirit of wine. It often contains variable proportions of propylic, butylic, and caproylic alcohols. See also VALERIANIC ACID.

Amylic alcohol is a "colorless liquid, with a penetrating and oppressive odor, and a burning taste. When pure its specific gravity is 0.818; boiling-point 279° F. Sparingly soluble in water, but soluble in all proportions in alcohol, ether, and essential oils. Exposed to the air in contact with platinum-black, it is slowly oxidized, yielding valerianic acid." Two allotropic varieties of amylic alcohol exist, one dextro-rotating a polarized ray.

ACETATE OF AMYL ($C_5H_{11}C_2H_3O_2$, or $Ay\overline{A}$).—To a small quantity of amylic alcohol in a test-tube add some acetate of potassium and a little sulphuric acid, and warm the mixture; the vapor of acetate of amyl is evolved, recognized by its odor, which is that of the jargonelle pear. If a condensing-tube be attached, the essence may be distilled over, washed by agitation with water to free it from alcohol, and separated by a pipette.

$$\underset{\text{Acetate of potassium.}}{K\overline{A}} + \underset{\text{Amylic alcohol.}}{AyHO} + \underset{\text{Sulphuric acid.}}{H_2SO_4} = \underset{\text{Acetate of amyl.}}{Ay\overline{A}} + \underset{\text{Acid sulphate of potassium.}}{KHSO_4} + \underset{\text{Water.}}{H_2O}$$

*Fruit Essences.*

Acetate of amyl, prepared with the proper equivalent proportions of constituents as indicated by the above equation, is largely manufactured for use as a flavoring agent by confectioners. Valerianate of amyl ($C_5H_{11}C_5H_9C_2$) is similarly used under the name of apple-oil. Butyrate of ethyl ($C_2H_5C_4H_7O_2$) closely resembles the odor and flavor of the pineapple; œnanthylate of ethyl ($C_2H_5C_7H_{13}O_2$) recalls greengage; pelargonate of ethyl ($C_2H_5C_9H_{17}O_2$) quince; suberate of ethyl

($Et_2C_9H_{12}O_4$) mulberry; sebacate of ethyl ($Et_2C_{10}H_{16}O_4$) melon. Hydride of salicyl ($C_7H_5O_2H$), or salicylous acid, is the essential oil of meadow-sweet (*Spiræa ulmaria*), and may be prepared artificially by the oxidation of salicin (*vide* p. 420). Salicylate of methyl ($CH_3HC_7H_4O_3$), or gaultheric acid (*Oleum Gaultheriæ*, U. S. P.), forms the chief part of the essential oil of wintergreen (*Gaultheria procumbens*), and may also be prepared artificially from salicin. Salicylic acid ($C_7H_6O_3$ or $H_2C_7H_4O_3$) can easily be obtained from the salicylate of methyl, but more cheaply from carbolic acid.

By mixing ethereal salts with each other and with essential oils in various proportions, the odor and flavor of nearly every fruit may be fairly imitated.

## Nitrite of Amyl.

Nitrite of Amyl (*Amyl Nitris*, B. P.) ($C_5H_{11}NO_2$).—For the preparation of this substance, Maisch recommends a modification of Balard's process. Amylic alcohol is first rectified until pure; that is, until its boiling-point is about 270° F. This alcohol, with about an equal bulk of nitric acid, is introduced into a capacious glass retort, and a moderate heat is applied and very gradually increased. As soon as the mixture approaches boiling, the fire is removed, and the reaction allowed to continue. If the application of the heat has been too rapid or too long continued, considerable frothing occurs, and the contents of the retort are apt to foam over. With a moderate and slowly increased heat, the reaction is less violent, and the temperature rises gradually after the removal of the fire and the beginning of boiling. As soon as the thermometer, inserted in the tubulus, rises above 212° F., the receiver is changed, the distillate now becoming more and more mixed with ethyl-amylic ether and nitrate of amyl, readily perceived by the change in odor.

The distillate obtained below 212° F. is agitated with an aqueous solution of hydrate or carbonate of potassium to remove free acids, and, after separation, the oily liquid is introduced into a clean retort and again slowly heated. The first portion coming over contains amylic aldehyd. When the very slowly increased heat has risen to 205° F., the receiver is again changed and the distillate now collected as nitrite of amyl, until the thermometer reaches 212° F.; when the distillation is stopped.

Nitrite of amyl is a yellowish ethereal liquid; sp. gr. of liquid 0.877, of vapor 4.03; boiling-point 205° F.; soluble in spirit of wine, insoluble in water; converted by fused caustic potash into valerianate of potassium.

---

## QUESTIONS AND EXERCISES.

780. Name the source of methylic alcohol.

781. What is "methylated spirit"?

782. Describe the method by which methylated spirit is detected in a tincture.

783. In what relation does formic acid stand to methylic alcohol?
784. How would you proceed to ascertain whether or not a specimen of sweet spirit of nitre had been made from methylated spirit?
785. Give details of the production of chloroform from alcohol, tracing the various steps by equations.
786. Is chloroform an ethylic compound? What is its probable constitution?
787. How is chloroform purified?
788. State the character of pure chloroform.
789. Describe the reactions that occur in the manufacture of chloral and chloral hydrate.
790. What is the nature of the action of alkalies on chloral hydrate?
791. Mention the characters of pure and impure chloral hydrate.
792. Whence is amylic alcohol obtained?
793. Has valerianic acid any chemical relation to amylic alcohol?
794. Mention the systematic names of several artificial fruit-essences.
795. What is the formula of nitrite of amyl, and how is it prepared?

---

## SALTS AND DERIVATIVES OF RADICALS OF OTHER SERIES THAN THE $C_nH_{2n+1}$.

What has been stated regarding radicals having the general formula $C_nH_{2n}+_1$ and their salts, may be applied to the radicals of other series.

The series $C_nH_{2n-7}$ includes *phenyl* ($C_6H_5$), the hydride of which ($C_6H_5H$, or PhH) is common *benzol* or *benzene* (B. P.), a colorless volatile liquid obtained from coal-tar, and a body which in constitution is the type of a very large class of aromatic organic bodies. *Toluol* ($C_7H_8$), or methyl-benzol, is one of the products of the distillation of balsam of Peru. The next homologue, *xylol* ($C_8H_{10}$), or dimethyl-benzol, is contained in crude wood-spirit and in coal-naphtha; it is a colorless fluid, having an odor faintly resembling that of benzol; boiling-point 282° F.; sp. gr. .866. Benzol is a powerful solvent of grease, and under the name of *Benzine Collas* was introduced by M. Collas, in 1848, for cleansing stuffs. By the action of strong nitric acid, benzol yields *nitrobenzol* ($C_6H_4(NO_2)H$), a liquid termed, from its odor, *artificial oil of bitter almonds*, or *essence of mirbane*. Its specific gravity is 1.20 to 1.29. The odor of this essence, however, is not exactly that of essential oil of almonds, and its composition is very different; so that it is not truly an artificial volatile oil. The natural oil has a sp. gr. of 1.04 to 1.07, and is a hydride of the negative radical* benzoyl ($C_7H_5OH$), a radical de-

---

* *Negative* basylous organic radicals are derived from the ordinary or *positive* basylous organic radicals by displacement of two atoms of hydrogen in the latter radicals by one atom of oxygen.

rived from the next higher homologue of phenyl by displacement of hydrogen by oxygen. Nitrobenzol yields aniline (*vide infra*), oil of bitter almonds does not (page 414). On this reaction the test for nitrobenzol is founded (p. 445). *Dinitrobenzol* ($C_6H_3(NO_2)_2H$) may also be obtained. The hydrate of phenyl ($C_6H_5HO$), or *phenic alcohol*, or *phenol*, is the *phenic acid* or *carbolic acid* ($H_2C_6H_4O$), of commerce (*Acidum Carbolicum*, U. S. P), a colorless crystalline substance, obtained from coal-tar oil by fractional distillation and subsequent purification. A crystalline hydrous acid ($H_2C_6H_4O,H_2O$) may also be obtained. At temperatures above 95° F. ordinary carbolic acid is an oily liquid. It is only slightly soluble in water, *Aqua Acidi Carbolici*, U. S. P., but readily dissolved by alcohol, ether, and glycerine (*Glyceritum Acidi Carbolici*, U. S. P.). In odor, taste, and solubility (and in appearance when liquefied by heat or by the addition of 5 per cent. of water) it resembles creasote, a wood-tar-product for which carbolic acid is often substituted. Besides hydrate of phenyl ($C_6H_5HO$) coal-tar oil contains *cresol*, *hydrate of cresyl*, or cresylic acid ($C_7H_7HO$); while wood-tar oil furnishes *guaiacol* ($C_7H_8O_2$)—also a product of the destructive distillation of guaiacum-resin—and *creasol* ($C_8H_{10}O_2$), or creasote. Certain coloring-matters may be obtained by the oxidation of carbolic acid; ammonia or, still better, phenyl-ammonia (aniline) mixed with it, and then a small quantity of solution of a hypochlorite, gives a blue liquid, a similar effect is produced on dipping a chip of deal into carbolic acid (or into creasote), then into hydrochloric acid, and afterwards exposing it to the air. By the following tests carbolic acid may be distinguished from creasote. The former boils only at 370° F., while the latter readily dries up at 212°. Carbolic acid does not affect a ray of polarized light; creasote twists it slightly to the right. Carbolic acid is either solid or may be solidified by cooling; creasote is not solidified by the cold produced by a mixture of hydrochloric acid and sulphate of sodium. Creasote from coal (impure or crude carbolic acid) gives a jelly when shaken with collodion; creasote from wood (*Creasotum* B. P. and U. S. P.) is unaffected by collodion (Rust). Coal-creasote is soluble in solution of potash and in the strongest solution of ammonia (Read), wood-creasote scarcely soluble. The coal product is soluble in a large volume of water (about 1 in 80, or 1 in 20 of boiling water), and a neutral solution of ferric chloride strikes a blue color with the liquid; wood-creasote is less soluble (*Aqua Creasoti*, U. S. P., is said to contain 1 in 129) and not colored blue by ferric chloride. An alcoholic solution of the coal-oil is colored brown by ferric chloride, a similar solution of true creasote green. According to Mr. Thomas Morson pure creasote is unaffected when mixed with an equal volume of commercial glycerine, while carbolic acid is miscible in all proportions, and will carry into solution even a considerable quantity of creasote. Carbolic acid is a powerful *antiseptic* (ἀντί, *anti*, against, and σήπω, *sepo*, I putrefy). In large doses it is poisonous, *antidotes* being a mixture of olive oil and castor oil, freely administered, or a mixture of slaked lime with about three times its weight of sugar rubbed together with a little water. *Acidum Carbolicum Impurum*, U. S. P., is the impure carbolic acid obtained from coal-tar by treating it first with an alkali, and then with an acid, and finally distilling

It is of a brown shade, and if at first colorless it becomes reddish-brown on pressure. It consists of carbolic and cresylic acids in variable proportion, together with impurities derived from coal-tar. 100 measures of warm water on being agitated with one measure of the liquid will dissolve out the two acids, and should not leave more than 30 per cent. by measure of impurity. Carbolic acid is soluble in oil of vitriol, *sulphocarbolic acid* ($HC_6H_5SO_4$) or *sulphophenic acid* being formed. On diluting and mixing with oxides, hydrates, or carbonates, *sulphocarbolates* are formed. The formula of *sulphocarbolate of sodium* is $NaC_6H_5SO_4$; *sulphocarbolate of zinc* $Zn(C_6H_5SO_4)_2H_2O$. Trinitro-carbolic acid ($C_6H_3(NO_2)_3O$) is formed on slowly dropping carbolic acid into fuming nitric acid; it is the yellow dye known as *carbazotic acid* or *picric acid;* most of the picrates are explosive by percussion. Both carbolic acid and benzol are secondary products, obtained in the manufacture of coal-gas; hence, indeed, the word *phenic* and thence *phenyl* (from φαίνω, *phaino*, I light, in allusion to the use of coal-gas). *Aniline*, or *phenylamine*, is a product of the action of nascent hydrogen on nitrobenzol.

$$C_6H_5NO_2 + 3H_2 = N\left\{\begin{matrix} C_6H_5 \\ H \\ H \end{matrix}\right. + 2H_2O,$$

Nitrobenzol. Hydrogen.

the substance whence, by oxidation, etc., aniline-red (magneta), -orange, -yellow, -green, -blue, -violet (mauve), and -black are produced. The process with zinc dust (*vide* Watt's Dict.) answers fairly well. 1 oz. of nitrobenzol gives a drachm or two of aniline. But action of iron filings and acetic acid on an alcohol solution answers better, 3 ozs. of nitrobenzol giving 1 oz. of aniline. To detect nitrobenzol add alcohol and hydrochloric acid and a little zinc; when action has ceased, liberate aniline by shaking with excess of potash; dissolve out the aniline by agitation with ether; to the decanted ethereal liquid add solution of chloride of lime; a violet color or tint results.

*Salicylic Acid* ($H_2C_7H_4O_3$).—The natural acid salicylate of methyl is described on page 442. Artificial salicylic acid is made as follows: Carbolic acid ($H_2C_6H_4O$), by acquiring the element of carbonic acid gas ($CO_2$) is converted into salicylic acid (Kolbe). The carbolic acid is first mixed with caustic soda in molecular proportions and dried. The resulting acid carbolate of sodium ($NaHC_6H_4O$), made hot, is saturated with carbonic gas, every pair of molecules of the carbolate thus affording one of regenerated carbolic acid, which distils off, and one of normal carbolate of sodium ($Na_2C_6H_4O$) which, absorbing the gas, becomes normal salicylate of sodium ($Na_2C_7H_4O_3$). The latter, by action of hydrochloric acid, furnishes salicylic acid, which may be purified by crystallization from alcohol. Salicylic acid, like carbolic acid, is a powerful antiseptic, but is free from the taste and smell of carbolic acid. It is only slightly soluble in water, but readily soluble in aqueous solutions of such alkali metal salts as borax, phosphate of sodium, or citrate of potassium. A similar antiseptic, *cresotic acid* ($C_8H_8O_3$), is similarly obtained from cresylic acid ($C_7H_8O$). Ferric

chloride strikes a violet coloration with both salicylic and cresotic acids. Both acids have antipyretic powers. The true salicylates of the alkali-metals, and probably, therefore, the cresotates are very feeble antiseptics. Salicylate of sodium is fairly soluble in water. Carbolic acid often containing cresylic acid, commercial salicylic acid may often contain cresotic acid. Salicylic acid is soluble in strong sulphuric acid, yielding *sulphosalicylic acid* ($HC_7H_5O_2SO_4$).

*Aloins.*—The aloes of pharmacy (*Aloe Barbadensis*, B. P., and *Aloe Socotrina*, B. P.) is an evaporated juice, doubtless much altered by the temperature to which it is subjected.

*Barbaloin.*—This substance, first obtained by T. and H. Smith, occurs in minute crystals in Barbadoes aloes. It is readily procured by making a decoction in water acidified with a little hydrochloric acid, after some hours pouring off the precipitated resin, and evaporating the liquid to a syrup. The aloin crystallizes out in a day or two. Barbaloin yields by the action of bromine and chlorine substitution compounds. Nitric acid dropped upon it produces a red color, which soon fades. Boiled for some time with strong nitric acid, barbaloin gives, together with oxalic and picric acids, a yellow substance, chrysammic acid, which furnishes beautiful red salts (Tilden). Anthracene ($C_{14}H_{10}$) has been obtained by deoxidation of barbaloin.

*Nataloin.*—This body was discovered by Flückiger in Natal aloes. It crystallizes readily in rectangular plates, either from spirit or from water. No bromine or chlorine substitution derivatives have yet been formed, but an acetyl compound has been analyzed (Tilden). Nataloin moistened with nitric acid gives a red coloration which does not fade. When boiled with nitric acid it yields no chrysammic, but only oxalic acid and picric acids.

*Socaloin*, or *Zanaloin.*—Histed and Flückiger have shown that Socotrine or Zanzibar aloes yields an aloin distinct from those just described. It forms tufted acicular prisms. Nitric acid scarcely alters the color of socaloin. Neither socaloin nor barbaloin affords any color when vapor from a glass rod moistened with nitric acid is brought near to a drop of oil of vitriol containing a minute fragment of the aloin, while nataloin gives rise to a blue coloration.

*Analysis.*—To Aloin, or powdered aloes—on a porcelain plate—add concentrated $HNO_3$. *No color* = *Socaloin.* *Crimson color* = Nataloin or Barbaloin. To another portion add concentrated $H_2SO_4$ and vapor of $N_2O_4$. *A blue color* = Nataloin. *No blue color* = Barbaloin.

*Formulæ.*—The reactions of these bodies seem to indicate that they are complex phenols. Phenol being the phenyl hydrate, $C_6H_5HO$, and cresol being methyl-phenol, $C_6H_4CH_3HO$, the aloins may possibly have a similar constitution—that is, they may be the hydrates of radicals in which part of the hydrogen is replaced by groups of atoms. Since Tilden made this suggestion, E. von Sommaruga and Egger (*Pharmacographia*) have arrived at the conclusion that the aloins form an homologous series, and that they have the composition indicated in the following formulæ:—

| | |
|---|---|
| Socaloin | $C_{15}H_{16}O_7$ |
| Nataloin | $C_{16}H_{18}O_7$ |
| Barbaloin | $C_{17}H_{20}O_7$ |

Tilden's more recent experiments indicate, however, that barbaloin ($C_{16}H_{18}O_7$) and socaloin ($C_{16}H_{18}O_7$) are isomeric in the anhydrous state, but that socaloin and its derivatives in the hydrous condition contain more water of crystallization than barbaloin. Nataloin ($C_{16}H_{18}O_7$) seems to be isomeric with the others, but is less soluble, and does not yield either chrysammic acid or chloro- or bromo-derivatives ($C_{16}H_{15}Cl_3O_7$; $C_{16}H_{15}Br_3O_7$). The acetyl derivative appears to have the formula $C_{16}H_{15}(C_2H_3O)_3O_7$.

*In the series* $C_nH_{2n-1}'$ we have the univalent radical *allyl* ($C_3H_5$) whose sulphide ($(C_3H_5)_2S$) is *essential oil of garlic*, and sulphocyanate ($C_3H_5CyS$), the chief part of the *essential oil of mustard*, the body to which mustard owes its power of inducing inflammatory action on the skin ("Mustard Poultice" and *Charta Sinapis*, B. P.). *Mustard* (*Sinapis*, B. P. and U. S. P.) is a powdered mixture of black, or rather reddish-brown, and white mustard-seeds. The white mustard-seed contains *sinalbin* ($C_{30}H_{44}N_2S_2O_{16}$), a glucoside which, in contact with the myrosin in an aqueous extract of mustard, yields some part of the acrid oil in the form of the sulphocyanide of a peculiar radical *acrinyl*.

$$\underset{\text{Sinalbin.}}{C_{30}H_{44}N_2S_2O_{16}} = \underset{\substack{\text{Sulphocyanate}\\\text{of acrinyl.}}}{C_7H_7OCNS} + \underset{\substack{\text{Acid sulphate}\\\text{of sinapine.}}}{C_{16}H_{24}O_5NSHSO_4} + \underset{\text{Glucose.}}{C_6H_{12}O_6}$$

The black contains the albumenoid ferment *myrosin*, resembling the *emulsin* of almonds (p. 414), and *myronate* of potassium or *sinigrin*. The latter is the body which, under the influence of the former, yields the chief part of the pungent oil.

$$\underset{\substack{\text{Myronate of}\\\text{potassium.}}}{KC_{10}H_{18}NS_2O_{10}} = \underset{\substack{\text{Acid sulphate}\\\text{of potassium.}}}{KHSO_4} + \underset{\substack{\text{Oil of}\\\text{mustard.}}}{C_3H_5CNS} + \underset{\text{Glucose.}}{C_6H_{12}O_6}$$

Crude oil of mustard often contains *cyanide of allyl*, $C_3H_5CN$.

In the Pharmacopœia of India the seed of *Sinapis juncea Rai*, or *Indian Mustard Plant*, is official in addition to that of *S. alba* and *S. nigra*. It is the common mustard of warm countries. It does not differ chemically from other mustard. Allyl compounds are also met with in several other cruciferous and liliaceous plants.

In the $C_nH_{2n}$ series occurs *ethylene* or *olefiant gas* ($C_2H_4$), the chief illuminating constituent of coal-gas (readily made on heating alcohol with twice its volume of strong sulphuric acid), a bivalent radical, the alcohol of which is *glycol* ($C_2H_4 2HO$). *Etherol* or *Ethereal Oil* (*Oleum Ethereum*, U. S. P., or *light oil of wine* ($C_{15}H_{30}$?), a hydrocarbon polymeric with olefiant gas, is one of the products of the action of excess of sulphuric acid on alcohol; its sp. gr. is 0.917. In $C_nH_{2n-1}'''$ series the trivalent hypothetical radical *glyceryl* ($C_3H_5$) is found, the hydrate of which ($C_3H_5 3HO$) is glycerine. The homologues of glycol are termed *glycols*, the homologues of glycerine

*glycerines.* It will be noticed that the chemical composition of the radicals glyceryl and allyl is identical ($C_3H_5$), but the former is trivalent and the latter univalent; hence they probably differ in physical constitution; they are isomeric, possibly polymeric with each other. From a glyceryl compound (glycerine), however, an allyl salt (iodide) can be produced. By distilling a mixture of glycerine and diniodide of phosphorus, iodide of allyl ($C_3H_5I$) is obtained, and on digesting this with sulphocyanate of potassium, the sulphocyanate of allyl, or *artificial oil of mustard*, results; identical with the natural oil.

GLYCERINE.—Glycerine, or Glyceric Alcohol ($C_3H_53HO$), is the hydrate of the basylous radical of most oils and fats, bodies which are mainly oleates, palmitates, and stearates of glyceryl (glycyl or propenyl). When these substances are heated with metallic hydrates (even with water—hydrate of hydrogen—at a temperature of 500° F. or 600° F.), double decomposition occurs, oleate, palmitate, or stearate of the metal is formed, and hydrate of glyceryl or glycerine. Hence glycerine is a by-product in the manufacture of soap, hard candles, and lead-plaster (*vide* Index).

*Properties.*—Glycerine is viscid when pure, specific gravity 1.28 (not below 1.25, B. P.), has a sweet taste, is soluble in water or alcohol in all proportions. It has remarkable powers as a solvent, is a valuable *antiseptic* even when diluted with 10 parts of water, and useful as an emollient. In vacuo it may be distilled unchanged, but under ordinary atmospheric pressure is decomposed by heat. When perfectly pure and anhydrous, glycerine, at a few degrees below the freezing point of water, sometimes solidifies to a mass of crystals.

*Tests.*—Heat one or two drops of glycerine in a test-tube, or with strong sulphuric acid, acid sulphate of potassium, or other salt powerfully absorbent of water; vapors of acrolein (from *acer*, sharp, and *oleum*, oil) are evolved, recognized by their powerfully irritating effects on the eyes and respiratory passages. If the glycerine be in solution in water, it must be evaporated as low as possible before applying this test.—Add a few drops of the fluid suspected to contain glycerine to a little powdered borax; stir well together; dip the looped end of a platinum wire into the mixture and expose to an air-gas flame: a deep green color is produced (Senier and Lowe).

The glycerine liberates boracic acid, and it is the latter which colors the flame. Ammoniacal salts, which similarly affect borax, must first be got rid of by boiling with solution of carbonate of sodium. Liquids containing much indefinite organic matter must sometimes be evaporated to dryness, the residue extracted by alcohol, and the latter tested for the glycerine. To detect traces liquids must be concentrated.

Besides glycerine itself (*Glycerinum*, B. P., *Glycerina*, U. S. P.), there are several official preparations of glycerine—solutions of carbolic, gallic, and tannic acids and borax in glycerine, and a sort of mucilage of starch in glycerine (*Glyceritum Acidi Carbolici*, *Glyceritum Acidi Gallici*, *Glyceritum Acidi Tannici*, *Glyceritum Picis Liquidæ*, and *Glyceritum Sodii Boratis*).

## QUESTIONS AND EXERCISES.

796. Give the formulæ of some compounds of radicals which have the general formula $C_nH_{2n-7}'$, $C_nH_{2n-1}'$, $C_nH_{2n-1}'''$ and $C_nH_{2n}''$,—*e. g.* benzol, essential oil of mustard, glycerine, and glycol.

797. State the difference in composition of natural and artificial oil of bitter almonds.

798. How is the so-called artificial oil of bitter almonds prepared?

799. What are the uses, composition, source, and properties of Carbolic Acid?

800. State the characters by which carbolic acid is distinguished from Creosote.

801. In what relation does carbolic acid stand to cresylic acid?

802. What is the general formula of sulphocarbolates?

803. Draw out an equation explanatory of the production of aniline.

804. Mention the chief properties of Glycerine.

805. What is the specific gravity of glycerine?

806. By what test is glycerine recognized?

807. Enumerate some official preparations in which glycerine is employed as a solvent.

## ALBUMENOID SUBSTANCES.

*Albumen.*—Agitate thoroughly, white of egg (*Albumen Ovi*, B. P.) with water, and strain or pour off the liquid from the flocculent membranous insoluble matter. One white to 4 ozs. of water forms the "Solution of Albumen," B. P.

*Test.*—Heat a portion of this solution of albumen to the boiling-point; the albumen becomes insoluble, separating in clots or coagula of characteristic appearance.

*Other Reactions.*—Add to small quantities of aqueous solution of albumen solutions of corrosive sublimate, nitrate of silver, sulphate of copper, acetate of lead, alum, perchloride of tin; the various salts not only coagulate but form insoluble compounds with albumen. Hence the value of an egg as a temporary antidote in cases of poisoning by many metallic salts,

its administration retarding the absorption of the poison until the stomach-pump or other measures can be applied. Sulphuric, nitric, and hydrochloric acids precipitate albumen; the coagulum is slowly redissolved by aid of heat, a brown, yellow, or purplish-red color being produced. Neither acetic, tartaric, nor organic acids, generally, except gallo-tannic, coagulate albumen. Alkalies prevent the precipitation of albumen.

*Yolk* or *Yelk of Egg* (*Ovi Vitellus*, B. P.) contains only 3 per cent. of albumen—the white 12½. The yolk also contains only 30 per cent. of yellow fat and 14 of casein. The whole egg (*Ovum*, U. S. P.) is also official.

Albumen is met with in large quantities in the serum of blood, in smaller quantity in chyle and lymph, and in the brain, kidneys, liver, muscles, and pancreas. It is not a normal constituent of saliva, gastric juice, bile, or mucus, but occurs in those secretions during inflammation. It is found in the urine and feces only under certain diseased states of the system.

The cause of the coagulation of albumen by heat has not yet been discovered.

Albumen has never been obtained sufficiently pure to admit of its composition being expressed by a trustworthy formula; Gerhardt regarded it as a sodium compound ($HNaC_{72}H_{110}N_{18}SO_{22}H_2O$).

Egg-albumen, and, to some extent, blood-albumen, is largely used by calico-printers as a vehicle for colors, serving also, when dry, as a glaze. Curriers prize egg-oil for softening leather.

Albumen coagulated by heat is said to be recoverable in a scarcely altered fluid condition by contact with a dilute aqueous solution of a very small proportion of pepsine.

### *Fibrin, Casein, Legumin.*

*Fibrin* is the chief constituent of the muscular tissue of animals. It occurs in solution in the blood; and its spontaneous solidification or coagulation is the cause of the *clotting* of blood shortly after being drawn from the body—a phenomenon which cannot at present be explained satisfactorily. Fibrin may be obtained by whipping fresh blood with a bundle of twigs, separating the adherent fibres, and washing in water till colorless. It may be dried or kept under spirit of wine.

Average Composition of Blood (in 1000 parts).
(Compiled by Kirkes.)

| | | | |
|---|---|---|---|
| | Water | | 784 |
| | Albumen | | 70 |
| | Fibrin | | 2.2 |
| | Red Corpuscles (dried) | | 130 |
| Fatty matters. | Cholesterin | 0.08 | 1.4 |
| | Cerebrin | 0.40 | |
| | Serolin | 0.02 | |
| | Oleic and margaric acids, Volatile and odorous fatty acid, Fat containing phosphorus | | |
| Inorganic salts. | Chloride of sodium | | 3.6 |
| | Chloride of potassium | | .35 |
| | Phosphate of sodium ($Na_3PO_4$) | | .2 |
| | Carbonate of sodium | | .82 |
| | Sulphate of sodium | | .28 |
| | Phosphates of calcium and magnesium | | .25 |
| | Oxide and phosphate of iron | | .50 |
| | Extractive matters, biliary-coloring-matter, gases, and accidental substances | | 6.4 |
| | | | 1000. |

Percentage proportion of the *chief* constituents of Blood.

| | |
|---|---|
| Water | 78.4 |
| Red corpuscles (solid residue) | 13.0 |
| Albumen of serum | 7.0 |
| Inorganic salts | .603 |
| Extractive, fatty, and other matters | .777 |
| Fibrin | .22 |
| | 100. |

*Casein* occurs in Cow's Milk (*Lac*, B. P.), to the extent of 3 per cent., dissolved by a trace of alkaline salt. Its solution does not spontaneously coagulate like that of fibrin, nor by heat like albumen; but acids cause its precipitation from milk in the form of a curd (cheese) containing the fat (butter)-globules previously suspended in the milk, a clear yellow liquid (or whey) remaining. *Curds and whey* are also produced on adding to milk a piece, or an infusion, of *rennet*, the salted and dried inner membrane of the fourth stomach of the calf. The exact action of rennet is not known. Respecting rennet, Soxhlet says, "60 to 80 grains of calf's stomach, steeped for five days in 1 litre of a 5 per cent. solution of common salt at ordinary temperatures, yield a solution of which 1 vol. will coagulate 10,000 vols. of new milk at a temperature of 95° F. in forty minutes. If the filtered solution is treated with 60 to 90 grains more of stomach, a solution of double strength is obtained; another repetition gives a solution three times the strength of the original one. To prevent decomposition, about 0.3 per cent. of thymol may be added to the

concentrated rennet extract solution. Possibly a slight taste due to this may be detected in the finest cheese, but for the same reason oil of cloves is much more objectionable. Boracic acid is on all accounts the best antiseptic to employ, and solutions to which it has been added may be kept in covered vessels for months. All extract solutions lose strength on keeping; during the first two months the solution may become thirty per cent. weaker, then the strength remains nearly constant for eight months in the case of a solution of 1 in 18,000. Alcohol is almost as good an antiseptic as boracic acid, if the solution be preserved in well-stoppered flasks."

*Average Composition of* 1000 *parts of Milk.*

| | Specific gravity. | Water. | Solid constituents. | Casein and extractive. | Sugar. | Butter. | Salts. |
|---|---|---|---|---|---|---|---|
| Woman . . . | 1.033 | 889 | 111 | 40 | 42 | 27 | 2 |
| Cow . . . . . | 1.034 | 864 | 136 | 55 | 37 | 36 | 7 |

Specific gravity alone, as taken by the form of hydrometer termed a *lactometer*, or even by more delicate means, is of little value as an indication of the richness of milk, the butter and the other solids exerting an influence in opposite directions. Good cow's milk affords from 10 to 12 per cent. by volume of cream, and 3 to 3½ per cent. of butter. The water of milk seldom varies more than from 87 to 88 per cent., and the solid constituents from 13 to 12. Indeed, excluding its butter, the milk of healthy cows is curiously regular in composition. The non-fatty solids in the mixed milk of a herd or dairy of healthy cows is almost a constant quantity, namely 9.3 per cent. A lower proportion of non-fatty solids in a sample of milk points to the addition of water. Thus, supposing that 100 grains of a specimen of milk evaporated to dryness, and all butter extracted from the residue by ether, yielded a non-fatty residue of 7.44 grains, the specimen would probably be four-fifths milk and one-fifth water. For if (by rule of three) 9.3 indicate 100, then 7.44 indicate 80. Occasionally, under exceptional circumstances, a sample of genuine milk might be slightly poorer than that from a healthy herd, and therefore, in England, for legal purposes, a standard of 9 per cent. by weight of non-fatty solids and 2.5 per cent. of butter-fat has been proposed. Only in the rare cases of milk containing an unusually large proportion of butter-fat would any milk yielding less than 9 per cent. of non-fatty solids be regarded as genuine. And, again, no milk would be considered genuine if it yielded less than 2.5 per cent. of fat, not even in the rare case of its containing an unusually large proportion of real non-fatty milk-solids. Half-starved cows might yield milk below these standards, but such milk could scarcely be considered to be genuine, or better fitted for food than milk watered *after* leaving the cow.

Under the microscope milk is seen to consist of minute corpuscles floating in a transparent medium. These corpuscles consist of fatty matter (butter), said to be contained in a filmy albumenoid envelope. The fat is fluid at the normal temperature of the animal, and remains so until the milk is well agitated by churning or otherwise, or until the milk is frozen.

*Legumin* or *vegetable casein* is found in most leguminous seeds, such as sweet and bitter almonds. Peas contain about 25 per cent. of legumin.

*Vegetable albumen* is contained in many plant-juices, and is deposited in flocculi on heating such liquids *Vegetable fibrin* is the name given by Liebig and Dumas to the portion of the gluten of wheat which is insoluble in alcohol and ether (*vide* p. 401).

*Albumenoid substances* are nearly identical in percentage composition. Albumen (and fibrin) contains 53.5 of carbon, 7 hydrogen, 15.5 of nitrogen, 22 of oxygen, 1.6 of sulphur, and .4 of phosphorus. Casein contains no phosphorus. These three bodies are often termed the *plastic elements of nutrition*, under the assumption that animals directly assimilate them in forming muscles, nerves, and other tissues, —starch, sugar, and similar matter forming the *respiratory* materials of food, because more immediately concerned in keeping up the temperature of the body by the combustion going on between them, and their products, and the oxygen of the air in the blood. The whole of the organic nitrogen in food must not, however, be regarded as representing true albumenoids, some existing as amidic and similar compounds, bodies having a simplicity of composition characteristic of the *products* of physiological action on food, rather than that complexity of composition characteristic of true nutrients. Albumenoids in decomposing yield much fatty as well as other substances. Possibly a portion, at least, of the *adipocire* (*adeps*, fat, *cera*, wax) or *corpse-fat*, characteristic of the remains of buried animals, is thus derived.

*Musk* (*Moschus*, B. P. and U. S. P.), "the inspissated and dried secretion from the preputial follicles of *Moschus moschiferus*" (the Musk-deer), is a mixture of albumenoid, fatty, and other animal matters with a volatile odorous substance of unknown composition.

---

## GELATIGENOUS SUBSTANCES.

These nitrogenous bodies differ chemically from the albumenoid in containing less carbon and sulphur and more nitrogen. They are contained in certain animal tissues, and on boiling with water yield a solution which has the remarkable property of solidifying to a jelly on cooling. The tendons, ligaments, bones, skin, and serous membranes afford *gelatine* proper; the cartilages give *chondrine*, which differs from gelatine in composition and in being precipitated by vegetable acids, alum, and the acetates of lead. The purest variety of gelatine is *isinglass*, B. P. (*Ichthyocolla*, U. S. P.), "the swimming-bladder or sound of various species of *Acipenser*, Linn., prepared and cut in fine shreds." Small quantities are more easily

disintegrated by a file than a knife. Fifty grains dissolved in 5 ounces of distilled water forms the official "Solution of Gelatine," B P. *Glue* is an impure variety of gelatine, made from the trimmings of hides; *size* is glue of inferior tenacity, prepared from the parings of parchment and thin skins. "Among the varieties of gelatine derived from different tissues and from the same sources at different ages, much diversity exists as to the firmness and other characters of the solid formed on the cooling of the solutions. The differences between isinglass, size, and glue, in these respects, are familiarly known, and afford good examples of the varieties called weak and strong, or low and high, gelatines. The differences are sometimes ascribed to the quantities of water combined in each case with the pure or anhydrous gelatine, part of which water seems to be intimately united with the gelatine; for no artificial addition of water to glue would give it the character of size, nor would any abstraction of water from isinglass or size convert it into the hard dry substance of glue. But such a change is effected in the gradual process of nutrition of the tissues; for, as a general rule, the tissues of an old animal yield a much firmer or stronger jelly than the corresponding parts of a young animal of the same species." (Kirkes's Physiology.)

Gelatine appears to unite chemically with a portion of the water in which it is soaked when used for culinary and manufacturing purposes, for a solution of glue in a hot anhydrous glycerine does not yield an ordinary jelly on cooling. From its solution in water, gelatine is precipitated by alcohol, corrosive sublimate, perchloride of platinum, and by tannic acid. Its aqueous solution is not, like that of albumen, coagulated by heat. By prolonged ebullition its gelatinizing power is destroyed.

---

## PEPSIN.

Pepsin (from πέπτω, *peptō*, I digest) is a nitrogenous substance existing in the gastric juice, and as a viscid matter in the *peptic* glands and on the walls of the stomachs of animals. To isolate it, the mucous membrane of the stomach (of the hog, sheep, or calf, killed fasting) is scraped, and macerated in cold water for twelve hours; the pepsin in the strained liquid is then precipitated by acetate of lead, the deposit washed once or twice by decantation, sulphuretted hydrogen passed through the mixture of the deposit with a little water to remove the whole of the lead, and the filtered liquid evaporated to dryness at a temperature not exceeding 105° F. Pepsin is a powerful promoter of digestion; its solution is hence frequently termed *artificial gastric juice*. As met with in pharmacy its strength varies greatly. It is often prepared by simply mixing with starch the thick liquid obtained on macerating the scraped stomach with water, and evaporating to dryness. (Vide *Pharmaceutical Journal*, 1865–66, p. 112, and 1871–72, pp. 785 and 843.) The official process (*Pepsin*, B. P.) simply consists in scraping the viscid pulp from the slightly washed inner surface of the stomach, and quickly evaporating it to dryness on glass or glazed earthenware at a temperature not

exceeding 100° F. The product is powdered. "A light yellowish-brown powder, having a faint, but not disagreeable odor, and a slightly saline taste, without any indication of putrescence. Very little soluble in water or spirit. Two grains of it with an ounce of distilled water, to which five minims of hydrochloric acid have been added, form a mixture in which 100 grains of hard-boiled white of egg, in thin shavings, will dissolve on their being digested together for about four hours at a temperature of 98°." The solvent or digestive action of pepsin on the albumenoids, etc., in the stomach results in a fluid termed *peptone*. It is thus that such food is prepared for conversion into blood. Artificial peptone is made by digesting blood-fibrin with pepsin in very weak hydrochloric acid. Peptone is not readily coagulated by heat, and it freely diffuses through membranes.

## PANCREATIN.

The pancreas (or "sweetbread") secretes a colorless fluid which contains 1½ to 2½ per cent. of an albumenoid substance, which has the power of converting starch into sugar and, especially, of emulsifying fat. It may be precipitated by chloride of sodium from an acidulated infusion of the pancreas. It is soluble in cold water. An extremely small proportion emulsifies a large volume of fat.

---

## QUESTIONS AND EXERCISES.

808. In what form is albumen familiar?
809. Name the chief test for albumen.
810. Why is the administration of albumen useful in cases of poisoning?
811. Mention the points of difference between yolk and white of egg.
812. From what source other than egg may albumen be obtained?
813. In what respect does fibrin differ from albumen?
814. Enumerate the chief constituents of blood?
815. How may fibrin be obtained from blood?
816. State the difference between casein, fibrin, and albumen.
817. What are the relations of cream, butter, curds and whey, and cheese, to milk?
818. Describe the microscopic appearance of blood and of milk.
819. How much cream should be obtained from good milk?
820. What is the percentage of water in genuine milk?
821. Name sources of vegetable casein and vegetable albumen.
822. Give the percentage of nitrogen in albumenoid substances.
823. Describe the chemical nature of musk.
824. In what lie the peculiarities of gelatine?
825. To what extent do isinglass, glue, and size differ?
826. Whence is pepsin obtained?
827. How is pepsin prepared?

# FATTY BODIES.

## SOAPS, SOLID FATS, FIXED OILS, VOLATILE OILS, CAMPHORS.

*General relations.*—Oils and fats are, apparently, almost as simple in constitution as ordinary inorganic salts. Just as acetate of potassium ($KC_2H_3O_2$) is regarded as a compound of potassium (K) with the characteristic elements of all acetates ($C_2H_3O_2$), so soft soap is considered to be a compound of potassium (K) with the elements characteristic of all oleates ($C_{18}H_{33}O_2$), and hence is chemically termed oleate of potassium ($KC_{18}H_{33}O_2$). Olive oil, from which soap is commonly prepared, is mainly oleate of the trivalent radical *glyceryl* ($C_3H_5$), the formula of pure fluid oil being $C_3H_53C_{18}H_{33}O_2$, and its name *oleine*. The formation of a soap, therefore, on bringing together oil and a moist oxide or hydrate, is a simple case of double decomposition or, rather, metathesis as seen already in connection with lead plaster (p. 213), or in the following equation relating to the formation of common hard soap:—

$$\underset{\text{Hydrate of sodium (caustic soda).}}{3NaHO} + \underset{\text{Oleate of glyceryl (vegetable oil).}}{C_3H_53C_{18}H_{33}O_2} = \underset{\text{Oleate of sodium (hard soap).}}{3NaC_{18}H_{33}O_2} + \underset{\text{Hydrate of glyceryl (glycerine).}}{C_3H_53HO}$$

Berthelot has succeeded in preparing oil artificially from oleic acid and glycerine; and it is said to be identical with the pure oleine of olive and of other fixed oils. Hard fats chiefly consist of *stearine*—that is, of tristearate of glyceryl ($C_3H_53C_{18}H_{35}O_2$). Mr. Wilson, of Price's Candle Company, obtains stearic and oleic acids and glycerine by simply passing steam, heated to 500° or 600° F., through melted fat. Both the glycerine and fat acids distil over in the current of steam, the glycerine dissolving in the condensed water, the fat-acids floating on the aqueous liquid.

The author finds that oleic acid readily combines with alkaloids and most of the metallic oxides or hydrates, forming oleates which are soluble in fats. In this way active medicines may be administered internally in conjunction with oils, or externally in the form of ointments.

As regards the conversion of oily substances into emulsions resembling the common natural emulsion, milk, Gregory states that three drachms of gum acacia in fine powder are necessary to emulsify one ounce of any of the volatile oils, and that a little less (about two drachms) will answer for the fixed oils and balsams. To this quantity of gum, four drachms and a half of water must be added (no more and no less). Either the water or the oil may be added first to the gum, but it is quickest to add the oil first, and well triturate before adding the water.

Soaps.—Olive oil boiled with solution of potash yields potassium soap, or *soft soap* (*Sapo Mollis*, B. P.); with soda, sodium soap, or *hard soap* (*Sapo Durus*, B. P. and U. S. P.); mixed with ammonia, an ammonium soap (*Linimentum Ammoniæ*, B. P. and U. S. P.); and with lime-water, calcium

soap (*Linimentum Calcis*, B. P. and—flaxseed oil—U. S. P.), —all oleates, chiefly of the respective basylous radicals. Their mode of formation is indicated in the foregoing equation. The alkali soaps are soluble in alcohol, the others insoluble. A green soap, much used on the continent of Europe, and, indeed, official in Germany (*Sapo Viridis*), is made by adding indigo to ordinary soft soap; the yellow color of the soap yielding with the indigo a greenish compound. The official characters of Hard Soap are: "grayish-white, dry, inodorous; horny and pulverizable when kept in dry warm air; easily moulded when heated; soluble in rectified spirit; not imparting an oily stain to paper; incinerated it yields an ash which does not deliquesce." And of Soft Soap: "yellowish-green, inodorous, of a gelatinous consistence; soluble in rectified spirit; not imparting an oily stain to paper; incinerated it yields an ash which is very deliquescent." Curd Soap (*Sapo Animalis*, B. P.) is a "soap made with soda and a purified animal fat, consisting principally of stearin." It will, of course, chiefly contain stearate of sodium. In pharmacy it is often advantageously employed instead of the "hard soap."

The hard soap met with in trade is made from all varieties of oil, the commoner kinds being simply the product of the evaporated mixture of oil and alkali; while the better sorts have been separated from alkaline impurities and the glycerine by the addition of common salt to the liquors, which causes the precipitation of the pure soap as a curd. Potash soap is not precipitable by salt.

*Bile*, the gall of the ox (*Bos taurus*, Linn.), freed from mucus by agitating with twice its bulk of rectified spirit (in which mucus is insoluble), filtering, and evaporating, yields the official Purified Ox-Bile (*Fel Bovinum Purificatum*, B. P.); the latter has a resinous appearance, but is chiefly composed of two crystalline substances having the constitution of a soap; the one is termed *taurocholate of sodium* ($NaC_{26}H_{42}NO_6S$), the other is *glycocholate*, or simply *cholate of sodium* ($NaC_{26}H_{44}NO_7S$). Both taurocholates and glycocholates are conjugated bodies readily yielding, the former *cholic* or *cholalic acid* ($HC_{24}H_{39}O_5$) and *taurine* ($C_2H_7NO_3S$), the latter cholalic acid and *glycocine* or *glycocoll* ($C_2H_5NO_2$), a soluble crystalline body having interesting physiological relations, inasmuch as it is obtainable from gelatine (hence the name glycocoll or sugar of gelatine, from γλυκὺς, *glucūs*, sweet, and κόλλα, *kolla*, glue) and hippuric acid. The presence of bile in a liquid, such as urine, is detected by Pettenkofer's test. The fluid is gradually mixed with half its bulk of strong sulphuric acid in a test-tube, rise of temperature being prevented by partial immersion of the tube in water. A small quantity of powdered white sugar is then introduced and well mixed with the acid liquid, and more sulphuric acid then poured in; as the temperature rises a reddish or violet coloration is produced. The cholacic acid liberated in the reaction furnishes the color.

SOLID FATS.—1. *Lard* (*Adeps Præparatus*, B. P. and U. S. P.) is the purified internal fat of the abdomen of the hog—the perfectly fresh *omentum* or *flare*, washed, melted, strained, and dried. 2. *Benzoated Lard* (*Adeps Benzoatus*, B. P.) is prepared lard heated over a water-bath with benzoin (10 grains per ounce), which communicates an agreeable odor and prevents or retards rancidity. Purified lard is a mixture of olcine and stearine; *margarine*, the margarate of glyceryl, was formerly supposed to be a constituent of lard and other soft fats, but is now regarded as a mere mixture of *palmitine* (the chief fat of palm oil) and stearine. 3. *Yellow Wax* (*Cera Flava*, B. P. and U. S. P.), the prepared honeycomb of the Hive-Bee, — and the same bleached by exposure to sunlight, 4, *White Wax* (*Cera Alba*, B. P. and U. S. P.),—according to Brodie, is chiefly a mixture of *Cerotic Acid* ($HC_{27}H_{53}O_2$), *Palmitate of Melissyl* ($C_{30}H_{61}C_{16}H_{31}O_2$), and about 5 per cent. of *Ceroleine*, the body to which the color, odor, and tenacity of wax are due. *Ceresine*, the purified native *ozokerite* (ὀζω, *ozō*, I smell; κηρός, *keros*, wax) of Galicia, is a solid hydrocarbon largely used as a substitute for beeswax, especially in Russia. It differs from beeswax chemically in being almost unattacked by warm oil of vitriol. 5. *Spermaceti* (*Cetaceum*, B. P. and U. S. P.) is the *palmitate of cethyl* ($C_{16}H_{33}C_{16}H_{31}O_2$), or *cetine;* when saponified it yields not glycerine, the hydrate of glyceryl ($C_3H_53HO$), but *ethal*, the hydrate of cetyl ($C_{16}H_{33}HO$); it is the solid crystalline fat accompanying *sperm* oil in the head of the spermaceti-whale. 6. *Suet*, the internal fat of the abdomen of the sheep, purified by melting and straining, forms the official *Prepared Suet* (*Sevum Præparatum*, B. P. and U. S. P.); it is almost exclusively composed of stearin ($C_3H_53C_{18}H_{35}O_2$). 7. Expressed oil of *nutmeg* (*Oleum Myristicæ Expressum*, B. P.), commonly but erroneously termed *Oil of Mace*, is a mixture of a little volatile oil with much yellow and white fat, the latter is *myristin* or myristate of glyceryl ($C_3H_33C_{14}H_{27}O_2$). 8. Oil of *theobroma*, or *Cacao-butter* (*Oleum Theobromæ*, B. P. and U. S. P.), chiefly theobromate of glyceryl (Kingzett), is a solid product of the roasted and roughly crushed seeds or *cocoa nibs* of the *Theobroma cacao*. They contain from one-fourth to one-half of this fat. These also furnish, when ground, *flake cocoa;* or when ground and much sweetened, *chocolate*; or, with farina and some sugar, *cocoa;* or with a portion of the butter extracted, "cocoatina," etc. 9. *Cocoa-nut* oil or butter, a soft fat largely contained in the edible portion of the nut of *Cocos nucifera*, or common cocoa-nut of the shops, a body containing glyceryl united with no less than six different univalent acidulous radicals, namely, the caproic ($C_6H_{11}O_2$), caprylic ($C_8H_{15}O_2$), rutic ($C_{10}H_{19}O_2$), lauric ($C_{12}H_{23}O_2$), myristic ($C_{14}H_{27}O_2$), and palmitic ($C_{16}H_{31}O_2$)—radicals which, like some from common resin, when united with sodium, form a soap differing from ordinary hard soap (oleate of sodium) by being tolerably soluble in a solution of chloride of sodium; hence the use of cocoa-nut oil and resin in making *marine soap*, a soap which, for the reason just indicated, readily yields a lather in sea-water. 10. *Kokum Butter*, Garcinia Oil, or Concrete Oil of Mangosteen, a whitish or yellowish-white fat obtained from

the seeds of *Garcinia indica* or *G. purpurea*, is composed of stearine, myristicine, and oleine. It is recognized officially in the Pharmacopœia of India (*Garciniæ purpureæ Oleum*).

FIXED OILS.—*Fixed* and *Volatile* oils are naturally distinguished by their behavior when heated; they also differ generally in chemical constitution, a fixed oil being, apparently, a combination of a basylous with an acidulous radical, while a volatile oil is more commonly a neutral hydrocarbon or the same oxidized.

*Drying and Non-drying Oils.*—Among fixed oils, most of which are oleate with a little palmitate and stearate of glyceryl, a few, such as, 1, *linseed oil* (*Oleum lini*, B. P. and U. S. P., contained in *Lini Semina*, B. P., *Linum*, U. S. P., or *Flaxseed*, the ground residue of which, after removal of much oil, is *linseed meal*, *Lini Farini*, B. P. and *Lini Farina*, U. S. P.), and, 2, *cod-liver oil* (*Oleum Morrhuæ*, B. P. and U. S. P.), and to some extent, *castor* and *croton*, are known as *drying oils*, from the readiness with which they absorb oxygen and become hardened to a resin. Linseed commonly contains 37 or 38 per cent. of oil; 25 to 27 per cent. is obtained by submitting the ground seeds to hydraulic pressure, 10 or 12 per cent. remaining in the residual *oil cake*. *Boiled oil* is linseed oil which has been boiled with oxide of lead. This treatment increases the already great tendency of linseed oil to resinify, forming *linoxyn*, $C_{32}H_{54}O_{11}$, on exposure to air. The drying oils appear to contain linoleine, an oily body distinct from oleine. Among the *non-drying oils* are the following: 3. *Almond* oil, indifferently yielded by the bitter (*Amygdala Amara*, B. P. and U. S. P.), or sweet seed (*Amygdala Dulcis*, B. P. and U. S. P.), to the extent of 45 and 50 per cent. respectively. 4. *Croton* oil (*Oleum Crotonis*, B. P., and *Oleum Tiglii*, U. S. P.). Geuther states that no such acid as crotonic is obtainable from croton oil, but acetic, butyric, valerianic, and higher members of the oleic series, together with *tiglic acid*, $HC_5H_7O_2$. 5. *Lycopodium*, a yellow powder composed of the spores of the common Club-Moss (*Lycopodium clavatum*), contains a large proportion of a very fluid fixed oil. 6. *Olive* oil (*Oleum Olivæ*, B. P. and U. S. P.), already noticed. 7. *Castor* oil (*Oleum Ricini*, B. P. and U. S. P.), chiefly a *ricinoleate of glyceryl* ($C_3H_5$-$3C_{18}H_{33}O_3$) or ricinoleine, a slightly oxidized oleine, soluble, unlike most fixed oils, in alcohol. Castor-oil seeds are stated, by Tuson, to contain an alkaloid, *ricinine*. It possesses no purgative property. 8. Oil of *male fern* (*Filix Mas*, B. P. and U. S. P.), a vermifuge obtained by exhausting the rhizome with ether and removing the ether by evaporation—a dark-colored oil containing a little volatile oil and resin, and officially termed an extract (*Extractum Filicis Liquidum*, B. P.). Its chief active constituent appears to be *filicic acid*, $C_{14}H_{18}O_5$. 9. Fixed oil of *mustard*, a bland, inodorous, yellow or amber oil, yielding, by saponification and action of sulphuric acid, glycerine, oleic acid, and *erucic acid* ($HC_{22}H_{41}O_2$) (Darby). 10. *Arachis* oil (*Oleum Arachis*, P. I.) is found to the extent of 40 or 50 per cent. in the seeds of the *Arachis hypogæa* (P. I.), the Ground-nut, or Earth-nut (so called because the pod of the herb in the growth of its stalk downward is forced beneath the surface of the ground and there ripens). It is chiefly olein, but contains hypogæin,

palmitin, and arachin. The oil is largely used in India in the place of olive oil, and is becoming much employed in Europe, especially for soap-making. 11. *Sesame* oil (Gingelly, Teal, or Benne Oil), from the seeds of *Sesamum indicum*, is also largely used in Europe. It has most of the characters of the best olive oil. 12. *Shark-liver oil*, from *Squalus carcharis* (*Oleum Squalæ*, P. I.), is used to some extent as a substitute for cod-liver oil in India.

| Acids. | Formulæ. Molec. Vol.=2. | Melting-point | | Boiling-point. | | Whence obtained. |
|---|---|---|---|---|---|---|
| | | ° F. | ° C. | ° F. | ° C. | |
| Formic ..... | $HC\ H\ O_2$ | 21 | —6 | 221 | 105.3 | Red ants; distillation of oxalic acid; and oxidation of amylaceous and other organic bodies. |
| Acetic ..... | $HC_2\ H_3\ O_2$ | 63 | 17 | 243 | 117 | Distillation of wood; oxidation of alcohol, etc. |
| Propionic .... | $HC_3\ H_5\ O_2$ | ..... | ..... | 284 | 140 | Fermentation of glycerine, etc. |
| Butyric ..... | $HC_4\ H_7\ O_2$ | below 0 | —20 | 314 | 157 | Butter; fermentation of lactic acid, etc. |
| Valerianic.... | $HC_5\ H_9\ O_2$ | " | " | 347 | 175 | Valerian root; oxidation of fousel oil. |
| Caproic..... | $HC_6\ H_{11}O_2$ | ..... | ..... | 392 | 200 | Butter. |
| Œnanthylic . | $HC_7\ H_{13}O_2$ | " | " | 298 ? | 148 ? | Castor oil by distillation, etc. |
| Caprylic ..... | $HC_8\ H_{15}O_2$ | 59 | 15 | 457 | 236 | Butter; cocoa-nut oil. |
| Pelargonic ... | $HC_9\ H_{17}O_2$ | ..... | ..... | 500 | 260 | Leaves of the geranium. |
| Rutic ..... | $HC_{10}H_{19}O_2$ | 86 | 30 | ..... | ..... | Butter; oil of rue by oxidation. |
| Lauric ..... | $HC_{12}H_{23}O_2$ | 110 | 43 | ..... | ..... | Cocoa-nut oil; berries of the bay tree. |
| Myristic..... | $HC_{14}H_{27}O_2$ | 129 | 54 | ..... | ..... | Nutmeg-butter; cocoa-nut oil, etc. |
| Palmitic ..... | $HC_{16}H_{31}O_2$ | 143.6 | 62 | ..... | ..... | Palm oil; butter; beeswax, etc. |
| Stearic ..... | $HC_{18}H_{35}O_2$ | 159 | 70.5 | ..... | ..... | Most solid animal fats. |
| Arachidic .... | $HC_{20}H_{39}O_2$ | 167 | 75 | ..... | ..... | Butter; oil of ground-nut. |
| Cerotic..... | $HC_{27}H_{53}O_2$ | 174 | 79 | ..... | ..... | Beeswax. |
| Melissic..... | $HC_{30}H_{59}O_2$ | 192 | 89 | ..... | ..... | Beeswax. |

By oxidation of oleic with nitric acid.

Their physical qualities and the formulæ of their acidulous radicals show that the fatty bodies are closely related, and indicate that the natural processes by which they are formed are probably as closely related. The above Table, from Miller's *Elements of Chemistry*, well shows the homology of the fat-acids, and gives their names, formulæ, melting-points, boiling-points, and natural and artificial sources. For further information respecting these acids and their isomers, the reader is referred to the Index, to works on "Organic Chemistry," to larger chemical works, or to Watt's *Dictionary of Chemistry*.

## QUESTIONS AND EXERCISES.

828. Give a sketch of the general chemistry of fixed oils, fats, and soaps.

829. What is the difference between Hard and Soft Soap?

830. Which soaps are official?

831. Name the source of lard, and state how "Prepared Lard" is obtained.

832. State the composition of Beeswax.

833. In what does Spermaceti differ from other solid fats?

834. Mention the chief constituent of Suet.

835. Whence is Cacao-Butter obtained?

836. Why is *marine soap* so called, and from what fatty matter is it almost exclusively prepared?

837. What do you understand by *drying* and *non-dryiny* oils?

838. In what respect does Castor Oil differ from other oils?

839. How is oil of male fern (*Ex. Filicis Liquidum*) prepared?

840. Mention the sources and formulæ of the following fat-acids: formic, acetic, proprionic, butyric, valerianic, caproic, œnanthylic, caprylic, pelargonic, and rutic.

---

VOLATILE OILS.—The *Volatile* or *Essential Oils* exist in various parts of plants, at first, probably, as mere combinations of carbon and hydrogen; but such *hydrocarbons* are prone to change when in contact with oxygen or moisture; hence these liquids, even when freshly obtained from the plants, and more especially as they occur in pharmacy, are usually mixtures of the liquid hydrocarbons or *elæoptens* (from ἔλαιον, *elaion*, oil, and ὄπτομαι, *optomai*, to see) pp. 461 to 467) with oxidized hydrocarbons, which are commonly solid or camphor-like bodies termed *stearoptens* (from στέαρ, *stear*, suet) (p. 467). The oils are also often associated with further oxidized bodies termed *resins* (p. 469). Of the hydrocarbons, those most commonly occurring are identical with or are isomers of that from oil of turpentine, and are known as *terpenes*. Tilden separates these terpenes into two groups: a turpentine group of true terpenes ($C_{10}H_{16}$) whose members boil at 156° C. to 160° C., yield a nitroso-derivative melting at 129° C., and form a solid crystalline hydrous terpene ($C_{10}H_{20}O_2,H_2O$); and an orange group ($C_{15}H_{24}$), polymerides of the true terpenes, whose members boil at 174° C. to 176°, yield a nitroso-derivative melting at 71° C., and form no solid hydrous compound. Kingzett's researches had previously pointed to a similar grouping, for he showed that turpentine and the other similar hydrocarbons ($C_{10}H_{16}$), and *cymene* ($C_{10}H_{14}$), the probable nucleus of these terpenes, when mixed with water and subjected to a current of air, yielded to the water peroxide of hydrogen (not ozone), camphor, and camphoric acid, the peroxide of hydrogen and camphoric acid possibly resulting from the action of water on a camphoric peroxide ($C_{10}H_{14}O_4$); while the second class ($C_{15}H_{24}$) yield no peroxide of hydro-

gen. The members of the $C_{10}H_{16}$ group are easily convertible into their polymers $C_{15}H_{24}$ and $C_{20}H_{32}$ by the action of heat, strong acids, etc. As regards either class, the members often differ from each other in their action on polarized light. Among the oxidized hydrocarbons occur several *aromatic aldehyds;* these, like ordinary aldehyds, form stable crystalline compounds with acid sulphate of sodium. On cooling a volatile oil, a stearopten often crystallizes out; or on distilling an oil, it remains in the retort, being less volatile than an elæopten. Volatile oils should obviously be preserved in well-closed bottles. Oxidation also proceeds more slowly in a cold than in a warm temperature.

*The process* by which volatile oils are usually obtained from herbs, flowers, fruit, or seeds, may be imitated on the small scale by placing the material (bruised cloves or caraways for instance) in a tubulated retort, adapting the retort to a Liebig's condenser, and passing steam, generated in a Florence flask, through a glass tube to the bottom of the retort. The steam in its passage through the substance will carry the oil over the neck of the retort into the condenser, and thence, liquefied and cooled, into the receiving vessel, where the particles of the oil will be found floating on the water. It may be collected by running off the distillate through a glass funnel having a stop-cock in the neck, or by letting the water from the condenser drop into an old test-tube which has a small hole in the bottom, or any similar tube placed in a larger vessel, the water and oil being subsequently run off separately from the tube as from a pipette. The water will in most cases be the ordinary official medicated water of the material operated on (*Aqua Aurantii Floris, Anethi, Carui, Cinnamomi, Fœniculi, Menthæ Piperitæ, Menthæ Viridis, Pimentæ, Rosæ*—from *Rosæ Centifoliæ Petala*, B. P. and U. S. P.—*Sambuci*). Volatile oils, like fixed oils, stain paper, but the stain of the former is not permanent like that of the latter. Oils of lemon and orange are sometimes obtained by mere pressure of the rind of the fruit.

The presence of alcohol in an essential oil may be detected and its quantity estimated by shaking with an equal bulk of pure glycerine. The latter dissolves the alcohol and is augmented in volume according to the amount of alcohol present (Boettger). For tests for the alcohol see page 425.

A large number of volatile oils are employed in medicine, either in the pure state, in the form of saturated aqueous solution (medicated waters), solution in spirit of wine, 1 in 5 (*Essentia Anisi*, and *Essentia Menthæ Piperitæ*, B. P.) and 1 in 50 (*Spiritus Cajuputi, Juniperi, Lavandulæ, Menthæ Piperitæ, Myristicæ, Rosmarini*), or as leading constituents in various barks, roots, leaves, etc. The strength of *Spiritus Anisi*, U. S. P., *Sp. Cinnamomi*, U. S. P.,

*Sp. Menthæ Piperitæ*, U. S. P., and *Sp. Menth. Viridis*, U. S. P., is 1 of oil to 15 of spirit of wine. *Perfumes* ("scents" or "essences," including "Lavender-Water" and "Eau de Cologne") are for the most part solutions of essential oils in spirit of wine, or spirituous infusions of materials containing essential oils. The following oils are, directly or indirectly, official in the Pharmacopœias: 1. Volatile oil of *Bitter Almond* (p. 414). 2. Oil of the fruits of *Ajwain* or *Omum*, *Carum Ajowan*, or *Ptychotis Ajowan* (*Fructus Ptychotis*, P. I.), contains *cymol* or *cymene* ($C_{10}H_{14}$) and a stearopten (*Ajwain-ka-phul*, flowers of ajwain) identical with *thymol*, $C_{10}H_{14}O$. 3. Oil of *Dill* (*Oleum Anethi*, B. P.), a pale, yellow, pungent, acrid liquid distilled from dill-fruit; it contains a hydrocarbon, *anethene* ($C_{10}H_{16}$), and an oxidized oil ($C_{10}H_{14}O$) identical with the carvol of oil of caraway (Gladstone). 4. Oil of *Aniseed* (*Oleum Anisi*, B. P.), a colorless or pale yellow liquid, of sweetish warm flavor, distilled in Europe from the Anise-fruit (*Pimpinella anisum*), and in China from the fruit of Star-Anise (*Illicium anisatum*); it is a mixture of a hydrocarbon isomeric with oil of turpentine and *anethol*, a stearopten ($C_{10}H_{12}$) which crystallizes out at low temperatures. 5. Oil of *Chamomile* (*Oleum Anthemidis*, B. P.), a bluish or, when old, yellow oil, of characteristic odor and taste, distilled from chamomile-flowers (*Anthemidis flores*, B. P.). The official variety (*Anthemis nobilis*) yields about 0.2 per cent. of an oil composed of a hydrocarbon ($C_{10}H_{16}$) and an oxidized portion ($C_{10}H_{16}O_2$) which, heated with potash, gives *angelate of potassium* ($KC_5H_7O_2$), whence is obtained *angelic acid* ($HC_5H_7O_2$). According to Demarcay the oil is a mixture of the angelates of butyl and amyl and similar bodies. The flowers of another variety (*Matricaria chamomilla*, U. S. P.) contain a stearopten ($C_{10}H_{16}O$) having the composition of laurel-camphor. 6. Oil of *Horseradish*-root (*Armoraciæ Radix*, B. P.) is, according to Hofmann, the sulphocyanate of butyl or tetryl ($C_4H_9CNS$); it is the chief active ingredient of *Spiritus Armoraciæ Compositus*, B. P. 7. Oil of Sweet-*Orange* peel (*Aurantii Dulcis Cortex*, U. S. P.) and Oil of Bitter-*Orange* rind (*Aurantii Cortex*, B. P.; *Aurantii Amara*, U. S. P.; *Aurantii Fructus*, B. P.), the flavoring constituent of the official syrup of the *peel* (*Syrupus Aurantii*, B. P.), and the oils of various species of *Citrus*,—namely, 8, *lemon* (*Oleum Limonis*, B. P. and U. S. P.), from Lemon Peel (*Limonis Cortex*, U. S. P.); 9, *lime;* 10, *bergamot* (*Oleum Bergamii*, U. S. P.); 11, *citron* and a variety of citron termed *cedra*,—resemble each other in composition, all containing *hesperidene*, a hydrocarbon ($C_{15}H_{24}$), and a small quantity of oxidized hydrocarbons ($C_{10}H_{10}O_5$, $C_{15}H_{10}O$, and (Wright and Piesse) $C_{20}H_{30}O_3$). Expressed lime essence also contains a soft resin. 12. Oil of *Neroli* or *Orange-Flower*, the aqueous solution of which is official in the forms of water (*Aqua Aurantii Floris*, B. P. and U. S. P.) and syrup (*Syrupus Aurantii Floris*, B. P. and U. S. P.), contains a fragrant hydrocarbon ($C_{10}H_{16}$), colorless when fresh, but becoming red on exposure to light, and an inodorous oxidized hydrocarbon. Strong acids, especially nitric, attack the oil in orange-flower water, coloring the fluid of a rose tint. 13. Oil of *Petit*

*Grain*, distilled from the leaves and shoots of the Orange tree, consists chiefly of a hydrocarbon apparently identical with that of oil of neroli. 14. The leaves of *Boldo* (*Peumus boldus*), a Chilian shrub (tonic and hepatic), yield two per cent. of essential oil (and, according to Bourgon and Verne, an alkaloid, *boldine*). 15. Oil of *Buchu-leaves* (*Buchu Folia*, B. P. and U. S. P.) consists of a hydrocarbon holding in solution a crystalline stearopten. Wayne has "more than once obtained from buchu a substance having the characters of salicylic acid." 16. Oil of *Cannabis Indica*, see p. 469. 17. Oil of (the lesser) *Cardomoms*, from seeds of the capsules (*Cardamomum*, B. P. and U. S. P.), is chiefly a hydrocarbon ($C_{10}H_{16}$) isomeric with oil of turpentine and a camphor resembling turpentine-camphor ($C_{10}H_{16}3H_2O$). 18. Oil of *Cajuput* (*Oleum Cajuputi*, B. P. and U. S. P.) is a mobile bluish liquid, consisting chiefly of *hydrous cajuputene* or *cajuputol* ($C_{10}H_{16}$,$H_2O$). The latter, repeatedly distilled from phosphoric anhydride, yields cajuputene itself ($C_{10}H_{16}$), which has the odor of hyacinths. Fresh cajuput oil has a green hue which is perhaps transient, for the color of the oil of trade is due to copper (Guibourt and Histed); certainly the green coloring matter of pure cajuput-oil is organic. 19. Oil of *Caraway*-fruit (*Carum*, U. S. P.) (*Oleum Carui*, B. P.; *Oleum Cari*, U. S. P.) is a mixture of *carvene* ($C_{10}H_{16}$) and *carvol* ($C_{10}H_{16}O$). 20. Oil of *Cloves* (*Oleum Caryophylli*, B. P. and U. S. P.) and of *Pimento* (*Oleum Pimentæ*, B. P. and U. S. P.), both heavier than water, contain a liquid hydrocarbon ($C_{15}H_{24}$), *eugenic acid* ($C_{10}H_{12}O_2$), a solid body, *eugenin*, isomeric with the eugenic acid, a second crystalline substance, *caryophyllin* ($C_{10}H_{16}O$), isomeric with common camphor, and a salicylic compound. 21. Oil of *Cascarilla*-bark (*Cascarillæ Cortex*, B. P. and U. S. P.) has not been fully examined. 22. Oil of *Cinnamon*-bark (*Cinnamomi Cortex*, B. P. and U. S. P.) and of *Cassia*-bark is mostly hydride of cinnamyl or cinnamic aldehyd ($C_9H_7OH$). Boiled with nitric acid it furnishes hydride of benzoyl ($C_7H_5OH$) and benzoic acid ($HC_7H_5O_2$), with chloride of lime yields benzoate of calcium ($Ca2C_7H_5O_2$), and with caustic potash gives cinnamate of potassium ($KC_9H_7O_2$). The specific gravity of oil of cinnamon (*Oleum Cinnamomi*, B. P. and U. S. P.) varies from 1.025 to 1.050. 23. Oil of *Citronella*, a *Grass Oil*, from *Andropogon nardus*, is chiefly composed of *citronellol* ($C_{10}H_{16}O$ and $C_{10}H_{18}O$, Wright), probably isomeric with the *absinthol* of wormwood (Gladstone). 24. Oil of *Capaiva* (*Oleum Copaibæ*, B. P. and U. S. P.) and, 25, of *Cubebs* (*Oleum Cubebæ*, B. P. and U. S. P.) are hydrocarbons, having the formulæ $C_{15}H_{24}$. This *cubebene* is sometimes associated with a camphor, *hydrous cubebene* ($C_{15}H_{24}$,$H_2O$). Oil of cubebs also contains a small quantity of a terpene ($C_{10}H_{16}$). 26. Oil of *Coriander* (*Coriandri Fructus*, B. P.; *Coriandrum*, U. S. P.; *Oleum Coriandri*, B. P.) seems to have the composition of hydrous oil of turpentine ($C_{10}H_{16}$-$H_2O$). 27. The fruits of *Cumin* or *Cummin* (*Cuminum cyminum*), an ingredient of many curry powders, contain about 3 per cent., and those of *Water Hemlock* or *Cowbane* (*Cicuta virosa*) about 1¼ per cent. of an essential oil composed of *Cymol* or *Cymene* ($C_{10}H_{14}$) and *Cuminol* ($C_{10}H_{12}O$). The latter is an aldehyd readily uniting with

alkaline bisulphites and by oxidation yielding *Cuminic acid* ($C_{10}H_{12}O_2$). Cymol also occurs in *Garden Thyme* (*Thymus vulgaris*). 28. *Eucalyptus globulus* furnishes an oil, the more volatile and chief portion of which is *eucalyptol* or *cymene* and a terpene ($C_{10}H_{14} + 2C_{10}H_{16}$), together with an oxidized portion, $C_{10}H_{14}O$ and $C_{10}H_{16}O$. 29. *Elecampane* root (*Inula Helenium*) by distillation with water yields solid volatile *helenin* ($C_6H_8O$), a camphor oil or *inulol* ($C_{10}H_{16}O$), and *inulic anhydride* ($C_{15}H_{22}O_3$). 30. Oil of *Fennel*-root (*Fœniculi Fructus*, B. P. and U. S. P.) differs in odor, but contains the same proximate constituents as oil of anise. 31. Oil of *Geranium*, or *Ginger Grass* oil, is from *Andropogon schœnanthus*, and contains *geraniol* ($C_{10}H_{18}O$). 32. *Grains of Paradise* (*Amomum melegueta*), Guinea Grains or Melegueta Pepper, *Semina Cardamomi Majoris*, contain essential oil ($C_{10}H_{16}$ and $C_{10}H_{16}O$) and a highly pungent resin. 33. Oil of *Juniper* (*Oleum Juniperi*, B. P. and U. S. P.), the active constituent of Juniper Tops and Berries (*Juniperus*, U. S. P.), contains a hydrocarbon ($C_{10}H_{16}$) which by contact with water yields a white crystalline hydrous compound ($C_{10}H_{16}H_2O$) and a polymeric hydrocarbon ($C_{20}H_{32}$). 34. Oil of *Lavender* (*Oleum Lavandulæ*, B. P. and U. S. P.) has not been satisfactorily examined. 35. Oil or butter or camphor of *Orris* (*Iris florentina*) is a soft solid lighter than water. Flückiger and Hanbury found it to be chiefly myristic acid associated with a little essential oil. 36. Oil of *Peppermint* (*Oleum Menthæ Piperitæ*, B. P. and U. S. P.) consists of a hydrocarbon, *menthene* ($C_{10}H_{18}$), different from that of most volatile oils, and hydrous menthene ($C_{10}H_{18}H_2O$) a crystalline stearopten. 37. Oil of *Spearmint* (*Oleum Menthæ Viridis*, B. P. and U. S. P.), the Common Mint of the kitchen garden, contains a liquid having the formula $C_{10}H_{20}O$, or $C_{10}H_{18}H_2O$; also, according to Gladstone, *menthol* ($C_{10}H_{14}O$) isomeric with carvol. 38. Oil of *Pennyroyal* (*Mentha pulegium*) contains, according to Kane, $C_{10}H_{16}O$. 39. Oil of *Nutmeg* (*Oleum Myristicæ*, B. P. and U. S. P.), and of the arillus of the nutmeg or mace (*Macis*, U. S. P.), is composed of a hydrocarbon, *myristicene* ($C_{10}H_{16}$), and *myristicol* ($C_{10}H_{16}O$)—Gladstone. 39*a*. Oil of *Origanum* (*Oleum origani*, U. S. P.), from *Origanum vulgare*, U. S. P., is of a bright yellow and has an odor somewhat like peppermint; it is a mixture of a liquid hydrocarbon and a camphor which is deposited after long standing. 40. Oil or *Otto* or *Attar* of *Cabbage-Rose Petals* (*Rosæ Centifoliæ Petala*, B. P. and U. S. P.; *Oleum Rosæ*, U. S. P.) gives the fragrance to Rose-water (*Aqua Rosæ*, B. P.). It resembles most other volatile oils in being composed of a hydrocarbon and an oxidized portion, but differs from all in this respect, that the hydrocarbon is solid and is destitute of odor, while the oxygenated constituent is liquid and the source of the perfume. According to Flückiger, the solid hydrocarbon ($C_{18}H_{16}$) yields succinic acid as the chief product of its oxidation by nitric acid, and in other respects affords evidence of belonging to the paraffin series of fats. 41. Oil of *Rosemary-tops* (*Oleum Rosmarini*, B. P. and U. S. P.) chiefly consists of a hydrocarbon ($C_{10}H_{16}$) resembling that from Myrtle, *Myrtus communis*, but also contains oxygenized oil and stearopten in variable proportions. 42. Oil of *Rue* (*Oleum Rutæ*,

B. P. and U. S. P.) contains a small quantity of hydrocarbon ($C_{10}H_{16}$) with some rutic aldehyd ($C_{10}H_{20}O$), but, according to Greville Williams, is chiefly euodic aldehyd ($C_{11}H_{22}O$), some lauric aldehyd ($C_{12}H_{24}O$) also being present. Gorup-Besanez and Grimm have obtained oil of rue ($C_{11}H_{22}O$) artificially as one of the products of the destructive distillation of acetate and caprate of calcium. 43. Oil of *Savin* (*Oleum Sabinæ*, B. P. and U. S. P.) contains several hydrocarbons isomeric with oil of turpentine (Tilden). 44. Oil of *Elder-flowers* (*Sambuci Flores*, B. P. and U. S. P.) occurs in very small quantity; it has a butyraceous consistence: it contains a hydrocarbon, *sambucene* ($C_{10}H_{16}$), and probably a camphor. 45. Oil of *Sandal Wood* has not been thoroughly examined. It occurs to the extent of about 1 per cent. in the fragrant white or yellow sandal wood of India, *Santalum album*, a small tree of the natural order Santalaceæ, and not to be confounded with the *Pterocarpus santalinus*, a tree of the natural order Leguminosæ, and furnishing the inodorous Red Sandal Wood or Red Sanders Wood of the dyer. 46. Oil of *Sassafras-root*, *Oleum Sassafras*, U. S. P., sp. gr. 1.094 (*Sassafras Radix*, B. P. and U. S. P.), yields *safren* ($C_{10}H_{16}$) and large quantities of a stearopten *safrol* or *sassafrol* ($C_{10}H_{10}O_2$). 47. Oil of *Mustard* (*Oleum Sinapis*, B. P.) is the sulphocyanate of allyl (p. 447). If adulterated with alcohol, its sp. gr. is below 1.015. 48. Oil of *Sweet Flag* (*Acorus calamus*) contains the hydrocarbon $C_{10}H_{16}$. (The rhizome also contains *Acorin*, a bitter glucoside.) 49. Oil of *Turpentine* (*Oleum Terebinthinæ*, B. P. and U. S. P.). Turpentine itself is really an oleo-resin of about the consistence of fresh honey. It flows naturally or by incision from the wood of most coniferous trees, larch (*Larix Europæa*) yielding *Venice turpentine*, *Abies balsameæ* furnishing *Canadian turpentine*, *Balsam of Fir*, or *Canada Balsam* (*Terebinthina Canadensis*, B. P. and U. S. P.), *Pistachia terebinthus*, the variety termed *Chian Turpentine*, and the *Pinus palustris*, *P. abies*, *P. pinaster*, and *P. tæda* affording the common, American Turpentine, *Terebinthus*, U. S. P. *Pinus maritima* gives the French or Bordeaux Turpentine, and *P. picea*, the old fragrant Strasburg Turpentine. By distillation turpentine is separated into *rosin* or *resin* (p. 469), which remains in the still, and *essential oil of turpentine*, often termed simply *turpentine*, *spirit of turpentine*, or "*turps*," which distils over. Mixed with alkali to saturate resinous acids, and redistilled, oil of turpentine furnishes about 80 per cent. of *rectified oil of turpentine*. Under the influence of heat, chemical agents, or both, pure oil of turpentine ($C_{10}H_{16}$) yields many derivatives of considerable chemical interest. The terpene of Bordeaux turpentine (terebenthene) rotates a ray of polarized light more than, and in the opposite direction to, the terpene of American turpentine. 50. Oil of common garden *Thyme* (*Thymus vulgaris*), *Oleum Thymi*, U. S. P., is composed of *cymene* or *cymol* ($C_{10}H_{14}$), *thymene* ($C_{10}H_{16}$), and *thymol* ($C_{10}H_{14}O$). Thymol is also contained in (*50a*) Oil of *Horsemint* (*Monarda*, U. S. P.). 51. Oil of *Turmeric* (*Curcuma longa*) resembles thymol. Thymol crystallizes out when oil of thyme or of ptychotis, etc., is kept at a low temperature for a day or two. It may also be obtained by shaking

the oils with caustic alkali and treating the separated alkaline liquid by an acid. It may be purified by distillation or crystallization from alcohol. It would seem that as an antiseptic thymol is far stronger than carbolic acid. 52. Oil of *Valerian-root* (*Valerinæ Radix*, B. P. and U. S. P.) (*Oleum Valerianæ*, U. S. P.) is a mixture of a hydrocarbon, *valerene* or *borneène* ($C_{10}H_{16}$), and *valerol* ($C_6H_{10}O$). Valerol slowly oxidizes to valerianic acid, known by its smell. A similar change occurs at once if oil of valerian be allowed to fall, drop by drop, on heated caustic potash: $C_6H_{10}O + 3KHO + H_2O = K_2CO_3 + KC_5H_9O_2 + 3H_2$. By the action of sulphuric acid on the valerianate of potassium thus produced, valerianic acid is obtained. 53. Oil of *Verbena*, *Lemon Grass* Oil, or *Indian Melissa* Oil, is obtained from *Andropogon citratus* (*Oleum Andropogi Citrati*, P. I.). 54. Oil of *Ginger* (*Zingiber*, B. P. and U. S. P.) has the composition of hydrous oil of turpentine. The pungent taste of ginger is due to a resin. 55. Wormseed (*Chenopodium*, U. S. P.) contains a volatile oil.

CAMPHORS.—In addition to the stearoptens or camphors already mentioned as being contained in or formed from volatile oils, there is one that is a common article of trade. It is obtained from the wood of *Camphora officinarum*, or Camphor-Laurel, in Japan (termed in Europe, Dutch camphor, because imported by the Dutch) and in China (known as Formosa camphor), by a rough process of distillation with water, and is resublimed in this country (*Camphora*, B. P. and U. S. P.). The formula of laurel-camphor is $C_{10}H_{16}O$. Bromine heated with camphor gives *monobrom camphor* ($C_{10}H_{15}BrO$) and hydrobromic acid. Monobrom-camphor is camphor in each molecule of which an atom of hydrogen has been displaced by one of bromine. Recrystallized, it occurs in white prisms. The essential oil (*Oleum Camphoræ*, U. S. P.), from which doubtless camphor is derived by oxidations, is easily obtained from the wood, and is occasionally met with in commerce under the name of *liquid camphor* or *camphor oil;* its formula is $C_{20}H_{32}O$; by exposure to air it becomes oxidized and deposits common camphor, $2C_{20}H_{32}O + O_2 = 4C_{10}H_{16}O$. Camphor distilled with phosphoric anhydride yields cymol, $C_{10}H_{14}$. There is another kind of camphor in European markets less common than laurel-camphor, but highly esteemed by the Chinese; it is obtained from the *Dryobalanops aromatica*, and denominated Sumatra or Borneo camphor. It differs slightly from laurel camphor in containing more hydrogen, its formula being $C_{10}H_{18}O$. It is accompanied in the tree by a volatile oil ($C_{10}H_{16}$) isomeric with oil of turpentine. This oil, *borneneè*, is also occasionally met with in trade under the name of *liquid camphor* or *camphor oil*, but differs from laurel-camphor oil in not depositing crystals on exposure to air.

Camphor is soluble to a slight extent in water (40 grains per gallon, Pooley). The official Camphor-water (*Aqua Camphoræ*, B. P. and U. S. P.), or *Camphor mixture*, is such a solution.

Common camphor, and many other of the camphors, oily hydrocarbons, and oxidized hydrocarbons, yield *camphoric acid* ($C_{10}H_{16}O_4$) and *camphoretic acid* ($C_{10}H_{14}O_7$) when attacked by oxidizing agents. Such reactions indicate natural relationships. Camphoric acid is a good antiseptic.

*Cantharidin* ($C_5H_6O_2$?), the active blistering principle of cantharides (*Cantharides*, B. P., *Cantharis*, U. S. P.) and other vesicating insects (such as *Mylatris cichorii* or *Telini Fly*, P. I., common in India), has most of the properties of a camphor or stearopten. It slowly crystallizes, from an alcoholic tincture of the beetles, in fusible, volatile, micaceous plates. The following process for the extraction of cantharidin is by Fumouze: Powdered cantharides are macerated with chloroform for twenty-four hours; and this treatment is repeated twice with fresh quantities of solvent, the residue having been well squeezed each time. The collected solutions are then distilled, and the dark-green residue treated with bisulphide of carbon, which dissolves fatty, resinous, and other matters and precipitates the cantharidin. The precipitate is thrown on a filter, washed with bisulphide of carbon, and recrystallized from chloroform. The same process, omitting the final recrystallization, may be used for the quantitative estimation of cantharidin in cantharides. The average quantity found is from four to five parts in one thousand. Cantharidin is readily soluble in warm glacial acetic acid (Tichborne) and still more readily in acetic ether.

Massing and Dragendorf consider cantharidin to be an anhydride ($C_5H_6O_2$), and that with the elements of water it forms *cantharidic acid* ($H_2C_5H_6O_3$). A cantharidate of potassium has the composition $KHC_5H_6O_3$. Piccard gives the vapor-density of cantharidin as about 6.5, and its formula $C_{10}H_{12}O_4$.

## QUESTIONS AND EXERCISES.

841. How do volatile oils usually differ chemically from fixed oils?

842. What are the general chemical characters of volatile oils?

843. Describe the usual process by which volatile oils are obtained.

844. Mention the differences in composition between the volatile oils of *Anthemis nobilis* and *Matricaria chamomilla*.

845. Give the systematic name of oil of horseradish.

846. State the general composition of the oil of lemon, lime, bergamot, citron, and cedra.

847. Name the constituents of oil of cloves.

848. In what respect does oil (or otto) of roses differ from other volatile oils?

849. To what class of substances do the constituents of oil of rue belong?

850. How does natural turpentine differ from the turpentine of trade?

851. With what object is commercial turpentine rectified?

852. How is camphor oil related to camphor?

853. In what respects do Borneo or Sumatra camphor and camphor oil differ from the corresponding products of Japan and China?

854. What is the nature of cantharidin?

## RESINOID SUBSTANCES.

### RESINS, OLEO-RESINS, GUM-RESINS, BALSAMS.

*Resins* occur in plants generally in association with volatile oils. They closely resemble camphors or stearoptens, but are not volatile, and differ from oils and fats mainly in being solid and brittle. *Oleo-resins* are mixtures of a resin and a volatile oil. *Gum-resins* are mixtures of a resin or oleo-resins and gum. *Balsams* are commonly described as resins or oleo-resins which yield benzoic or cinnamic acid; but oleo-resins containing neither of these acids are often termed balsams, *e. g.*, balsam of copaiva and Canada balsam. A physico-chemical method for the identification of the chief resins, gum-resins, and balsams, will be found in the *Pharmaceutical Journal* for Nov. 17, 1877.

RESINS.—1. *Resin*, *rosin*, or *colophony* (*Resina*, B. P. and U. S. P.) is the type of this class. Its source is the oleo-resin or true turpentine of the conifers, a body which by distillation yields spirit of turpentine and a residuum of rosin. "Brown" and "White" rosin are met with in trade. The former is the residue of American, the latter of Bordeaux turpentine (from *Pinus Abies*, etc., and *Pinus Maritima*, respectively). The chief constituents of brown resin are *pinic acid* ($HC_{20}H_{29}O_2$) and *sylvic acid*, identical in composition, but differiug in properties (*vide* Isomerism), the former being soluble and the latter insoluble in cold spirit of wine. White resin or "galipot" is chiefly *pimaric acid*, also isomeric with pinic acid. Pinic acid heated yields *colophonic* or *colopholic acid*. Among the products of the destructive distillation of resin, Tichborne has recently found "*colophonic hydrate*" ($C_{19}H_{22}O_3, H_2O$), a white inodorous crystalline substance, and by depriving this of water has obtained white crystalline *colophonine* ($C_{19}H_{22}O_3$). Resin is soluble in oil of turpentine. Contact with sulphuric acid immediately colors it strongly red. It is a constituent of eight of the fourteen Plasters (*Emplastra*) of the British Pharmacopœia. 2. *Arnicin* ($C_{20}H_{30}O_4$), the chief acrid if not the only active principle of Arnica (*Arnicæ Radix*, B. P., *Arnicæ Flores*, U. S. P.), is a resin. 3. *Cannabin*, said to be the active principle of *Indian Hemp* (*Cannabis Indica*, B. P., *Extractum Cannabis*, U. S. P.), was obtained in 1846 by T. and H. Smith, and is a resin. Personne in 1857 isolated a volatile oil, also said to possess much medicinal activity, consisting of *cannabene* ($C_{18}H_{20}$) and a solid crystalline "*hydride of cannabene*" ($C_{18}H_{22}$). Preobraschensky states that the active principle is nicotine. 4. *Capsicum-fruit* contains resin (p. 470). 5. *Castorin*, a resinous matter, is the name given to the chief constituent of *Castor* (*Castoreum*, B. P. and U. S. P.), the dried preputial follicles and included secretion of the Beaver (*Castor Fiber*). 6. *Copal*. The best copal is the exuded resin of trees of extinct forests, and is found beneath the surface of the ground in the neighborhood of existing trees. It appears to be a mixture of acids, but its character is still obscure. 7. *Dragon's Blood*, a crimson-red resin found as an exudation on the mature fruits of a Rotang or Rattan Palm

(*Calamus draco*). It is said to be a definite substance, and to have the formula $C_{20}H_{20}O_4$. 8. *Ergotin* is a very active resinoid constituent of *Ergot* (*Ergota*, B. P. and U. S. P.), or the "sclerotium (compact mycelium or spawn) of *Clariceps purpurea*, produced within the paleæ of the common rye, *Secale cereale*." According to Wenzell, ergot contains two alkaloids, *ecboline* and *ergotine*, to the former of which, he says, the activity of ergot is due. Blumberg considers these alkaloids to be identical. Tanret states that an unstable alkaloid termed *ergotinine* occurs in ergot and to the extent of 1 per 1000, and that it is accompanied by a camphor. Dragendorff and Podwissotzky assert that ergot owes most of its activity to *sclerotic* or *sclerotinic acid*, present to the extent of about 4 per cent. 9. *Guaiacum-resin* is a mixture of several substances (p. 418). 10. *Jalap-resin* (p. 419). 11. *Kousso* (*Cusso*, B. P.) yields yellow crystals of a resinoid body readily soluble in alkaline liquids, *kosin* or *koussin* ($C_{31}H_{38}O_{10}$). 12. *Mastic* (*Mastiche*, B. P. and U. S. P.) is a resinous exudation obtained by incision from the stem of the Mastic or Lentisk tree. Nine-tenths of Mastic is *mastichic acid* ($C_{20}H_{31}O_2$), a resin soluble in alcohol; the remaining tenth, *masticin* ($C_{70}H_{81}O$), a tenacious elastic resin. 13. *Mezereon*, the dried bark (*Mezerei Cortex*, B. P. and U. S. P.) of *Daphne mezereum*, Mezereon, and *Daphne laureola*, Spurge Laurel, owes its acridity to a resin. 14. *Pepper* contains a resin (p. 397). 15. *Burgundy Pitch* (*Pix Burgundica*, B. P. and U. S. P.) is the melted and strained exudation from the stem of the Spruce Fir, *Abies Excelsa*. The term Burgundy is a misnomer, the resin never having been collected at or near Burgundy; Finland, and to a smaller extent Baden, and Austria being the countries whence it is derived. Its constituents closely resemble those of common Resin. It is often adulterated and imitated by a mixture of resin with palm-oil, water, etc., from which it may be readily distinguished by its duller yellow color, highly aromatic odor, greater solubility in alcohol, and almost complete solubility in twice its weight of glacial acetic acid (Hanbury). 16. *Podophyllum resin* (p. 395). 17. *Pyrethrin* is the name of the acrid resinous active principle of the root of *Anacyclus pyrethrum* or *Pellitory-root* (*Pyrethri Radix*, B. P.). According to Buckheim the action of alkalies breaks it up into piperidine and pyrethric acid. (The crystalline poisonous principle obtained by Bellesme from *Pyrethrum carneum*, the powder of which is the well-known "insecticide," has not yet been analyzed.) 18. The resins of Rhubarb have already been alluded to in connection with Crysophanic acid (p. 337). 19. *Rottlerin* is the name given by Anderson to a crystalline resin from Kamala (*Kamala*, B. P.), the minute glands that cover the capsules of *Rottlera tinctoria:* to this, and, apparently, allied resins, Kamala owes its activity as an anthelmintic.

Oleo-resins.—1. *Capsicin*, a term suggestive of a definite chemical substance, is a name somewhat unhappily accorded to an indefinite substance, an oleo-resin, obtained by digesting the alcoholic extract of Capsicum fruit (*Capsici Fructus*, B. P.) in ether and evaporating the clear ethereal fluid to dryness. Besides volatile oil and resin, capsicum fruits contain much fatty matter which Thresh states is

chiefly free palmitic acid. (See also *Capsicine* and *Capsaicin*, p. 395.) 2. *Copaiva* (*Copaiba*, B. P. and U. S. P.) is a mixture of essential oil ($C_{15}H_{24}$), with two or more per cent. of brown soft resin, and 30 to 60 of a yellow dark resin termed *Copaivic acid* ($C_{20}H_{30}O_2$). Copaiva, containing about equal parts of this acid and of the oil, heated with a fourth of its weight of the official carbonate of magnesium yields a transparent fluid, owing to the formation of copaivate of magnesium and solution of this soap in the essential oil. With an equal weight of the carbonate enough soap is produced to take up the whole of the essential oil, and form a mass capable of being rolled into pills. A much smaller quantity of calcined magnesia, as might be expected, effects the same result; but more time, often several days, is required before complete reaction is effected. Quicklime has a similar effect. Perhaps carbonate reacts more quickly because of its fine state of division and admixture of hydrate—in which case hydrates of calcium and magnesium may be expected to act better than the calcined preparations, and in much smaller quantity than carbonate of magnesium. Copaiva, unlike, 3, *Wood-oil* or *Gurjun Balsam* (*Dipterocarpi Balsamum*, P. I.), a similar oleo-resin from the *Dipterocarpus turbinatus* (*D. Lævis*, P. I.), does not become gelatinous when heated to 270° F. Copaiva is often slightly fluorescent; Gurjun balsam is highly fluorescent. The stated analogy of Gurjun balsam to copaiva is borne out by its chemical composition; for by distillation it yields about 40 per cent. of an essential oil identical in composition with oil of copaiva, the non-volatile portion being resinous. The adulteration of copaiva with fixed oil is best detected by heating 20 or 30 drops in a capsule until all essential oil has evaporated. Turpentine is betrayed by its odor during this evaporation. The residue, copaiva resin, is brittle if pure, and more or less sticky or soft if fixed oil is present. The limit of brittleness is stated, by Siebold, to be reached when one per cent. of oil has been added to the copaiva, that amount preventing the residue being reduced to a fine powder. 4. *Elemi* (*Elemi*, B. P.) is an exudation from a tree growing in the Philippine Islands. It consists of volatile oil ($C_{10}H_{16}$) with 80 or more per cent. of two resins, the one ($C_{20}H_{32}O_2$) soluble in cold alcohol, the other *amyrin*, $(C_5H_8)_5H_2O$, almost insoluble, associated with *amyric acid*, $(C_5H_8)_{11}O_4$ (Buri). It also contains small quantities of two crystalline bodies soluble in water, *Bryoidin*, $(C_5H_8)_4 3H_2O$, and *Breidin* (Flückiger). The *Icacin* of Stenhouse and Groves is either identical with amyrin or perhaps has the formula $(C_5H_8)_9H_2O$. All these bodies are probably hydrous terpenes. 5. *Wood-Tar* (*Pix Liquida*, B. P. and U. S. P.) is a mixture of several resinoid and oily bodies (amongst others Creasote, p. 444) obtained by destructive distillation from the wood of *Pinus sylvestris* and other pines. When heated it yields a terebinthinate oil and a residue of *pitch*. 6. *Turpentines*. These oleo-resins have been mentioned in connection with oil of turpentine, their volatile, and resin, their fixed constituent. 7. *Common Frankincense* (*Thus Americanum*, B. P.) is the concrete turpentine of *Pinus tæda*. 8. *Canada Balsam* (*Terebinthina Canadensis*, B. P.), largely gathered in the province of Quebec, is the turpentine or oleo-resin of the Balm of Gilead Fir

(*Abies balsamea*). 9. *Sumbul-root* (*Sumbul Radix*, B. P.) seems to owe its stimulating property to two oleo-resins, one soluble in ether, the other in alcohol. 10. *Oleo-resin of Lupulin* (U. S. P.) is an ethereal extract of the yellow powder (*Lupulina*, U. S. P.) attached to the small nuts at the base of the scales which form the aggregate fruit of the Hop (*Humulus Lupulus*). It contains essential oil of hop (*Valerol*, $C_6H_{10}O$) and oxidized oil or resin, bitter extract containing the hop-bitter, lupulinic acid ($C_{32}H_{50}O_7$), and tannic acid. *Oleoresinæ Capsici*, *Cubebæ*, *Piperis*, and *Zingiberis* are also official in the United States Pharmacopœia. 11. *Pix Canadensis*, U. S. P., is the concrete juice of *Abies Canadensis*.

GUM-RESINS.—1. *Ammoniacum* (*Ammoniacum*, B. P. and U. S. P.) is an exudation from the *Dorema Ammoniacum*. It contains nearly 20 per cent. of gum, a little volatile oil, and about 70 of resin ($C_{40}H_{50}O_9$—Johnston). 2. *Asafœtida* (*Asafœtida*, B. P.) is a gum-resin obtained, by incision, from the living root of *Narthex asafœtida*. It contains from 50 to 70 per cent. of a resin, which is partly *ferulaic acid* ($C_{10}H_{10}O_4$), 25 to 30 per cent. of gum (about two-thirds arabin, one-third bassorin, p. 114), and 3 to 5 per cent. of volatile oil. 3. *Euphorbium*, an old drug which is an emetic and purgative resin. It contains an amorphous active resin ($C_{30}H_{48}O_4$), crystalline *euphorbon* ($C_{26}H_{44}O_2$), and mucilage (Flückiger). 4. The ordinary or Siam *Gamboge* (*Cambogia*, B. P. and U. S. P.) of European trade is obtained from the *Garcinia morella*; the gamboge of India (*Cambogia Indica vel Mysoriensis*, P. I.) from *G. pictoria*. When of best quality it contains about 20 per cent. of a gum, and 80 to 75 per cent. of a resin termed *gambogic acid* ($C_{20}H_{22}O_4$). 5. *Galbanum* (*Galbanum*, B. P. and U. S. P.) contains from 20 to 25 per cent. of gum, about 65 per cent. of resin ($C_{40}H_{54}O_7$), and 3 or 4 per cent. of volatile oil. Galbanum heated for some time to 212° F. with hydrochloric acid, the liquid separated and shaken with ether or chloroform, and the latter evaporated, yields somewhat less than 1 per cent. of colorless acicular crystals of *umbelliferone* ($C_9H_6O_3$).* 6. *Myrrh* (*Myrrha*, B. P. and U. S. P.), an exudation from the stem of *Bal-*

* "Umbelliferone is soluble in water; its solution exhibits, especially on addition of an alkali, a brilliant blue fluorescence which is destroyed by an acid. If a small fragment of galbanum is immersed in water, no fluorescence is observed, but it is immediately produced by a drop of ammonia. The same phenomenon takes place with asafœtida, and a slight degree with ammoniacum; it is probably due to traces of umbelliferone pre-existing in those drugs. Umbelliferone is also produced from many other aromatic umbelliferous plants, as *Angelica*, *Levisticum*, and *Meum*, when their respective resins are submitted to dry distillation; also from the resin of *Daphne mezereum*. The fluorescence of umbelliferone may be beautifully shown by dipping some bibulous paper into water which has stood for an hour or two on lumps of galbanum, and drying it. A strip of this paper placed in a test-tube of water with a drop of ammonia will give a superb blue solution, instantly losing its color on the addition of a drop of hydrochloric acid." (Flückiger and Hanbury.)

*samodendron myrrha*, contains about half its weight of soluble gum (arabinoid), 10 per cent. of insoluble gum (probably bassorin), 2½ of volatile oil, and about 25 per cent. of resin (myrrhic acid). 7. *Olibanum* (P. I.), *Thus musculum*, or *Arabian Frankincense* (from various species of *Boswellia*), is about one-third gum and nearly two-thirds resin ($C_{40}H_{30}O_6$) with a little hydrocarbon ($C_{10}H_{16}$) and oxidized hydrocarbon volatile oils. It has always been an important ingredient of *incense*—myrrh, storax, benzoin, and such fragrant combustible resinous substances being other constituents. 8. *Scammony* (p. 421).

Gum-resins need only be finely powdered and rubbed in a mortar with water to yield a medical *emulsion*, in which the fine particles of resin are held in suspension by the aqueous solution of gum.

BALSAMS.—1. *Benzoin* (*Benzoinum*, B. P. and U. S. P.) is obtained from incisions of the bark of *Styrax benzoin*. It contains 12 to 15 per cent. of benzoic acid (p. 335), about 50 per cent. of a resin (α) soluble in ether, 25 to 30 per cent. of a resin (β) soluble in alcohol only, and 3 to 4 per cent. of a resin (γ) soluble in solution of carbonate of sodium. The α resin is considered to be a compound of the β ($C_{40}H_{46}O_8$) and the γ ($C_{39}H_{40}O_5$). The balsams of Peru, Tolu, and Storax differ from benzoin in containing cinnamic (p. 462) in place of benzoic acid; hence they yield, by oxidation, hydride of benzoyl (oil of bitter almonds). 2. *Balsam of Peru* (*Balsamum Peruvianum*, B. P. and U. S. P.), from the *Myroxylon Pereiræ*, is a mixture of oily matter with about one-quarter or one-third of resinous matter, and 6 per cent. of cinnamic acid. The oil by fractional distillation in an atmosphere of carbonic acid gas and under diminished pressure, furnishes *benzylic alcohol* ($C_7H_7HO$), *benzoate of benzyl* ($C_7H_7C_7H_5O_2$), and *cinnamate of benzyl* ($C_7H_7C_9H_7O_2$) or *cinnamein* (Kraut). By action of alcoholic solution of potash it yields benzoate and cinnamate of potassium, and benzylic alcohol; also *cinnamic alcohol* ($C_9H_9HO$), otherwise known as *peruvine* or *styrone*; it also often holds in solution *metacinnamein* or *styracin* ($C_{18}H_{16}O_2$), isomeric with hydride of cinnamyl ($C_9H_7OH$). The resin of balsam of Peru seems to result from the action of moisture on the oil. The constituents of *Vanilla*, U. S. P., the prepared unripe pods of a plant, somewhat resemble those of Balsam of Peru. The crystals often found on vanilla are a weak acid substance having the formula $C_{16}H_{16}O_6$ (Carles). 3. *Balsam of Tolu* (*Balsamum Tolutanum*, B. P. and U. S. P.) is an exudation from incisions in the bark of *Myroxylon toluifera*; in composition it closely resembles balsam of Peru, but is more susceptible of resinification. It contains benzoate and cinnamate of benzyl, cinnamic acid, a little benzoic acid (Busse), and about 1 per cent. of a volatile hydrocarbon, *tolen*, $C_{10}H_{16}$. Old hard balsam of tolu is a convenient source of cinnamic acid, which is extracted by the same process as that by which benzoic acid may be obtained from benzoin, namely, ebullition with alkali, filtration, and precipitation by hydrochloric acid. 4. *Storax* is an oleo-resin obtained from the *Liquidambar orientale*. It contains a volatile oil termed *styrol cinnamene* or *cinnamol* ($C_8H_8$)—which possibly (Berthelot) is condensed acetylene, $4C_2H_2$—cinnamic acid, styracin or

cinnamate of cinnamyl ($C_9H_9C_9H_7O_2$), and a soft and a hard resin. Styrol differs from similar hydrocarbons in being converted into a polymeric solid termed *metastyrol* or *draconyl* on the mere application of a temperature of about 400° F. For medicinal use, storax (*Styrax Præparatus*, B. P. and U. S. P.) is purified by solution in alcohol, filtration, and removal of the alcohol by distillation.

### *Caoutchouc* or *India rubber*, and *Gutta Percha*.

Caoutchouc is the hardened juice of *Hevea* (*Siphonia*) *Brasiliensis*, *Castilloa elastica*, *Urceola elastica*, *Ficus elastica*, and other plants (Collins). Heated moderately with sulphur it takes up 2 or 3 per cent., and forms *vulcanized India-rubber*; at a higher temperature a hard horny product, termed *ebonite* or *vulcanite*, results. *Gutta Percha* (B. P. and U. S. P.) is the concrete drop or juice of the *percha* (Malay) tree, the *Isonandra gutta*, and of other Sapotaceous plants. It is soluble in chloroform (*Liquor Gutta-percha*, U. S. P.), benzoyl, and essential oils. White gutta percha is obtained by precipitating a solution of the ordinary gutta percha in chloroform by alcohol, washing the precipitate with alcohol, and finally boiling in water and moulding into the desired form while still hot. The official solution of Gutta Percha (*Liquor Gutta Percha*, B. P) is made by digesting *thin* slices of gutta percha in 12 parts by weight of chloroform, and then "fining" by shaking with 1 part of carbonate of lead and setting aside till the fluid is clear.

These two elastic substances, in the pure state, are hydrocarbons ($xC_5H_4$), usually slightly oxidized.

---

## QUESTIONS AND EXERCISES.

855. How do resins occur in nature? Distinguish between resins and camphors. Mention the points of difference of resins, oleo-resins, gum-resins, and balsams.

856. Name the source and chief constituents of common Resin or Rosin.

857. Enumerate some official articles of which the active constituents are resins.

858. Give the chief distinguishing characters of Burgundy Pitch.

859. What is the average proportion of oil and of resin in the so-called Balsam of Copaiva?

860. Explain the effect of carbonate of magnesium, magnesia, and lime on copaiva.

861. State the nature of Wood-tar.

862. Why do Ammoniacum, Asafœtida, Gamboge, Galbanum, Myrrh, and similar substances, give an emulsion by mere trituration with water?

863. In what respect does Benzoin differ from the Balsams of Peru, Tolu, and Storax?

864. What is the chemical nature of India-rubber and Gutta Percha?

865. How is India-rubber *vulcanized* and converted into *ebonite* or *vulcanite*?

---

## COLORING-MATTERS.

The animal, vegetable, and mineral kingdoms abound in substances or pigments which powerfully decompose light, absorbing certain of its constituent colors, and reflecting some others. Thus, for example, most leaves contain a body termed chlorophyll, which has the property of absorbing red light and reflecting green; these reflected rays entering the eye of an observer, and striking on the retina (the expanded extremity of the optic nerve), always communicate the same impression to the brain; in popular language the leaf is said to be green. Art has richly supplemented the number of such natural coloring-matters.

YELLOW.—1. *Chrome-yellow* occurs in more than a dozen shades (see *Lead, chromate of*). 2. *Fustic* or *yellow wood* is the wood of the *Rhus cotinus*. 3. *Gamboge* (see *Gamboge*). 4. *Ochre* is met with of many tints, under the names of *yellow ochre*, *gold yellow*, *gold earth*, or *ochre*, *yellow sienna*, *Chinese yellow*. It is chiefly a mixture of oxyhydrates of iron with alumina and lime. 5. *Orpiment* is a sulphide of arsenicum ($As_2S_3$). 6. *Persian berries* or *Avignon grains* contain a yellow principle termed *rhamnin* and other crystalline bodies. They are the product of the *Rhamnus infectorius*. 7. *Purree* or *Indian yellow* is said by Stenhouse to owe its color to *purrate* or *euxanthate of magnesium* ($MgC_{42}H_{34}O_{22}$). 8. *Quercitron* is the bark of *Quercus tinctoria*, U. S. P., or Black Oak. It contains the yellow glucoside, *quercitrin* ($C_{18}H_{16}O_{10},H_2O$). 9. *Rhubarb* (see *Chrysophanic acid*, p. 337). 10. *Saffron* (*Crocus*, B. P. and U. S. P.), the dried stigma and part of the style of *Crocus sativus*, yields *saffranin* or *polychroite*, an orange-red glucoside, which by the action of dilute acids and by other means breaks up as shown in the following equation, yielding red *crocin* (Weiss).

$$\underset{\text{Polychroite.}}{C_{48}H_{60}O_{18}} + \underset{\text{Water.}}{H_2O} = 2(\underset{\text{Crocin.}}{C_{16}H_{18}O_8}) + \underset{\text{Vol. oil of saffron.}}{C_{10}H_{14}O} + \underset{\text{Sugar.}}{C_6H_{12}O_6}.$$

11. *Turmeric*, the rhizome of *Curcuma longa*, owes its yellow color to *curcumin*, a resinous matter, the formula of which is said by Daube to be $C_{10}H_{10}O_3$, and by Iwanof $C_4H_4O$. Possibly two yellow pigments are present. 12. *Weld* (*Reseda luteola*) contains a durable yellow matter termed *luteolin* ($C_{20}H_{14}O_8$). 13. *Picric* or *carbazotic acid* (p. 445) is a very powerful yellow dye. 14. Dried and powdered carrots yield to bisulphide of carbon a yellow coloring-matter, "carrotin," which is obtained on evaporating the solvent. It is said to be used in coloring butter.

RED.—1. *Alkanet*, the root of *Alkanna tinctoria*, Tausch, *Anchusa tinctoria*, Desf., yields *anchusin* ($C_{26}H_{40}O_8$), a resinoid matter

soluble in oils and fats. 2. *Annato*, *Arnatto*, or *Arnotto*, a paste prepared from the seeds of *Bixa orellana*, contains *bixin*, an orange-red, and *orellin*, a yellow principle. 3. *Brazil-wood* (*Cæsalpinia brasiliensis*) furnishes *brezilin*, the basis of several lakes. *Sapan-wood* and *Camwood* probably contain the same substance. 4. *Cinnabar*, Chinese red, Vermilion, or Paris red is mercuric sulphide. 5. *Chrome-red* is an oxychromate of lead. 6. *Cochineal* (p. 337). 7. *Madder*, the root of *Rubia tinctorum*, powdered and treated with sulphuric acid and acidulated water to effect the removal of earthy and other inert matters, furnishes a residual powder termed *garancin*. Garancin yields to pure water *alizarin* ($C_{14}H_8O_4$, $3H_2O$), the red, neutral, crystallizable coloring-matter of madder. Alizarin does not exist ready formed in the plant, but is derived, by fermentation, from rubian, a yellowish resinoid substance. Alizarin is now produced artificially from *anthracene*, one of the solid constituents of coal-tar. 8. *Mulberry-juice* (*Mori Succus*, B. P.) contains a violet-red coloring matter which has not been chemically examined. 9. *Red lead* (p. 212). 10. *Red oxide of iron*, of shades varying from light to brown-red, is found native. The common names of it are Armenian bole, Berlin-red, colcothar, English red, red ochre, burnt ochre, red earth, terra di sienna, mineral purple, stone red, and Indian red. 11. *Red Sanders-wood* or *Red Sandal-wood* (*Pterocarpi Lignum*, B. P. and U. S. P.), the billets and chips of *Pterocarpus santalinus*, owes its color to *santalin* ($C_{14}H_{12}O_4$), a crystalline resinoid matter. 12. *Red-Poppy Petals* (*Rhœdos Petula*, B. P.), from the *papaver rhœas*, contain a red coloring principle which has not yet been isolated in a state of purity. The author has sought for morphia in large quantities of the petals, but could not find a trace of that alkaloid. 13. *Red-Rose Petals* (*Rosæ Gallicæ Petala*, B. P. and U. S. P.), and those of the *Cabbage Rose* (*Rosæ Centifoliæ Petalæ*, B. P), also yield a red substance, which has not been analyzed. 14. *Safflower*, *Dyer's Saffron* or *Bastard Saffron*, the florets of *Carthamus tinctorius*, contains an unimportant yellow dye, and 5 per cent. *carthamin* ($C_{14}H_{16}O_7$), an uncrystallizable red dye, the pigment of the old *pink saucers*. Carthamin seems to possess acid characters, and (like silicic acid and other substances) to be soluble in water for a certain time after liberation from its alkaline solution; for fabrics are dyed with safflower by immersion in a bath made of an infusion in dilute alkali neutralized by citric acid immediately before use, the carthamin probably penetrating the cells and vessels of the fibres in a soluble form, there becoming insoluble and imprisoned and thus giving permanent color to the wool, silk, or other material. Mixed with French chalk, carthamin is used as a cosmetic under the name of *vegetable rouge—carmine* being *animal rouge*, and *red oxide of iron mineral rouge*. 15. *Lac-dye* is a cheap form of cochineal, and is also yielded by the species of *Coccus*, whose resinous excretion constitutes *shellac*. 16. *Logwood* (*Hæmatoxyli Lignum*, B. P.) contains a yellow substance, *hæmatoxylin* ($C_{16}H_{14}O_6$, $H_2O$ or $3H_2O$), which, under the influence of air and alkali, assumes an intense red color. Under the influence of ammonia and air it yields greenish-violet iridescent scales

of *hæmatin* ($C_{16}H_{12}O_6,3H_2O$). 17. *Red enamel* colors, for glass-staining and ceramic operations, are produced either by cuprous silicate or purple of Cassius (p. 246).

BLUE.—1. *Cobalt oxide* precipitated in combination or admixture with alumina or phosphate of calcium forms *Thénard's blue*, *cobalt-blue*, *Hoffner's blue*, and *cobaltic ultramarine*. 2. *Smalt*, *Saxony blue*, or *King's blue*, is rough cobalt glass in fine powder (p. 234). 3. *Copper-blue*, *mountain-blue*, and *English* or *Hambro' blue* are carbonates or oxycarbonates of copper. 4. *Indigo* (p. 290). 5. *Litmus*, *lichen-blue*, *turnsole*, *orchil* or *archil*, and *cudbear* are products of the action of air and alkalies on certain colorless principles, as *orcin* ($C_7H_8O_2$), derived from different species of lichen—*Roccella*, *Variolaria*, and *Lecanora*. 6. *Prussian blue* (p. 340) and *Turnbull's blue* (p. 341), the ferro- and ferridcyanides of iron, are met with under the names of *Erlangen*, *Louisa*, *Saxon*, *Paris*, or *Berlin blue*. 7. *Ultramarine* is made on a large scale by roasting a mixture of fine white clay, carbonate of sodium, sulphur, and charcoal. Its constitution is not well made out. Acids decompose it, sulphuretted hydrogen escaping.

GREEN.—1. *Cupro-arsenical* green pigments (p. 177). 2. *Chlorophyll*, *Leaf-green*, or *Chromule*. A method of extracting chlorophyll is given under "Extracts," *vide* Index. It is resinoid, soluble in alcohol and ether, insoluble in water, and, according to Frémy, consists of a blue substance, *phyllocyanin* ($C_{34}H_{68}N_4O_{17}$?), and a yellow, *phylloxanthin;* the yellow tints in fading autumnal leaves, he says, are due to the latter principle, the former being the first to fade. Chlorophyll would probably well repay extended investigation. 3. *Sap-green*, *buckthorn-*, *vegetable-*, or *bladder-green* is obtained by evaporating to dryness a mixture of lime and the juice (*Rhamni Succus*, B. P.) of the berries of the Buckthorn (*Rhamnus catharticus*). It is soluble in water, slightly in alcohol, and insoluble in ether and oils. 4. *Green ultramarine* is made by a process similar to that for blue ultramarine. 5. Mixtures of blue and yellow pigments and dyes are common sources of green colors. 6. Glass and earthenware are colored green by oxide of chromium and black oxide of copper.

BROWN.—1. *Umber*, *Sienna*, or *Chestnut-brown* is found native. By heat it is darkened in tint, and is then known as *burnt umber*. It is a mixture of oxide of iron, silica, and alumina. 2. *Sepia* is a dried fluid from the ink-bag of cuttle-fishes (*Sepiadæ*); by its ejection into adjacent water the animal obtains opportunity of escape from enemies. 3. *Catechu* (p. 357) furnishes a brown coloring-matter.

BLACK.—1. *Blacklead* (p. 30), *bone-black* (p. 111), or *ivory-black*, and *lampblack*, the latter a deposited soot from incomplete combustion of resin and tar, are varieties of carbon. 2. *Burnt sugar* or *caramel* (p. 412). 3. *Indian ink* is usually a dried mixture of fine lampblack and size, or thin glue. 4. *Black ink* is essentially tannates and gallates of iron suspended in water containing a little gum in solution 5. *Printer's ink* is well-boiled linseed or other oil, mixed with good lampblack, vermilion, or other pigment. 6. *Black dyes* are of the same nature as ink. 7. The *pigmentum nigrum* of

black feathers, such as those of the common rook, of dark hair, and probably also of the skin of the negro, seems to be due to the black substance which remains undissolved when black feathers are digested for some time in dilute sulphuric acid. It is said to have the formula $C_{18}H_{16}N_2O_8$ (Hodgkinson and Sorby).

White Pigments.—1. *Chalk* or *Whiting* (p. 105). 2. *French chalk*, *steatite*, or *soapstone*, a silicate of magnesium. 3. *Heavy white* (p. 102). 4. *Pearl-white* (p. 252). 5. *Plaster of Paris* (p. 114). 6. *Starch* (p. 401). 7. *White-lead* (p. 210). 8. *Zinc-white* (p. 133). 9. Oxides of tin and zinc and phosphate of calcium are employed for giving a white opacity to glass.

Aniline Colors. *Coal-tar colors.*—Within the last ten years nearly every shade of color seen in the animal or vegetable kingdoms has been successfully imitated by certain dyes and pigments primarily derived from a mineral, coal. Coal distilled for gas furnishes tar or gas-tar. Coal-tar contains some aniline; but especially it contains a liquid convertible into aniline, namely, benzol ($C_6H_5H$), first discovered by Faraday in compressed oil-gas. From aniline, by oxidation, Runge obtained the violet color-reaction, the body producing which Perkin afterwards studied and isolated, and manufactured under the name of *mauve*. *Aniline-red* (*fuchsine*, *magenta*, or *rosaniline*), *aniline-yellow*, *aniline-green*, *aniline-blue*, and, in short, aniline-dyes, lakes, and pigments of every hue of the rainbow, are now common articles of trade. Their application has revolutionized the arts of the dyer and color printer.

## QUESTIONS AND EXERCISES.

866. Explain the production of color, by various natural and artificial pigments.

867. Mention the chief yellow coloring-matters, and describe their chemical nature.

868. What is annatto?

869. Name the colorific constituent of madder. Can it be made artificially?

870. State the source of Litmus.

871. Distinguish between Prussian blue and Turnbull's blue, and state how they are manufactured.

872. How is blue ultramarine obtained? How is it affected by acids?

873. Describe the chemical nature of the coloring principle of green leaves.

874. By what agents is glass colored green?

875. Whence is sepia obtained?

876. Describe the chemistry of black ink.

877. Write a few sentences on *aniline colors*.

## QUALITATIVE ANALYSIS OF SUBSTANCES HAVING UNKNOWN PROPERTIES.

Substances are presented to the analyst in one of the three forms in which all matter exists—namely, solid, liquid, or gaseous—and they may contain animal or vegetable as well as mineral matter.

The method of analysis in the case of *solid mineral bodies* has been described on pp. 369 to 376.

*Solid animal or vegetable substances* (or mixtures of these with mineral bodies) may be indefinite and beyond the grasp of chemistry, or definite and quite within the range of proximate qualitative organic analysis. The presence of such substances is indicated in the preliminary examination of a solid (pp. 369 to 371) by charring and other characters. If no charring occurs, and no volatile liquid is expelled by heat, the absence of such matter is indicated. But if organic matter is present, an endeavor is made to ascertain its precise character. The analyst's knowledge of the history of the substance, or the circumstances under which it comes into his hands, will probably afford a clue to its nature, and enable him to search directly for its proximate constituents. If no such information is at hand, the action of solvents may be employed as likely to afford indication of the general, if not of the precise nature of the substance. Water, alcohol, ether, chloroform, bisulphide of carbon, each both hot and cold, may in turn be agitated with the substance, the mixture filtered, a portion of the filtrate evaporated, at first partially, setting the product aside, and afterwards to dryness, and any deposit or residue examined with and without the aid of a microscope. Other portions of the filtrate may be treated with acids, alkalies, and solutions of such metallic salts as are commonly used as group-tests for acidulous radicals (p. 366). The action of alkalies, as well as acids weak and strong, hot and cold, may also be tried on the solid substance itself, and colors, odors, and, in short, any effect whatever, duly noted. A portion of the substance should also be burnt in an open porcelain crucible until no carbon remains, and the ash, if any, examined; its amount and nature may afford information leading to the identification of the substance.

The foregoing experiments having been carefully performed, and all results entered in the note-book, a little reflection will

possibly lead to the recognition, or may suggest further direct experiments or confirmatory tests, or will, at least, have pointed to the absence of 90 or 95 per cent. of all possible substances, and thus have restricted the area of inquiry to narrow limits. The success attainable in qualitative proximate organic analysis by the medical or pharmaceutical student will of course largely depend on the thoroughness with which the operator has prosecuted his study of practical chemistry generally, but it also will be considerably affected by the extent to which he has cultivated the art of observation, and the opportunities he has had of acquiring a knowledge of the appearances, uses, and common properties of definite chemical substances, and of articles of food, drink, and medicine. The most successful of several good analysts will be he who has most common sense and most experience.

The pharmaceutical student, who has probably already had some years of experience in pharmacy, occupies an unusually favorable position for prosecuting the proximate analysis of organic and inorganic substances, or, at all events, of that large proportion of such bodies met with in the domain of hygiene and pharmacy. Many substances he will indentify at sight, or by aid of a lens, or after applying some simple physical or chemical test. Nor should he find much difficulty, after reaching the present point of practical study, in deciding whether the solid substance under examination belongs to the class of organic acids, organic salts of metallic radicals, alkaloids, salts of alkaloids, amylaceous matter, gums, saccharine substances, glucosides, albumenoid matters, fats, soaps, resins, color-matters, etc. For instance, the pharmaceutical student will find less difficulty than the general student in successfully analyzing a substance occurring in "scales," because he has experience of the appearances of compounds commonly produced in that form, and because, even if the appearance is new to him, he knows what kind of substances most readily lend themselves to production in that form. While the general student is testing generally, and proceeding cautiously, or searching for general information in books of reference, the pharmaceutical or medical student has incinerated some of the material, noticed whether or not the ash is red (iron) and strongly alkaline (potassium), treated more of the material with an alkali (for ammonium), added excess of ammonia, and examined the precipitate (for cinchonia or quinia), or shaken up the alkaline liquid successively with ether and chloroform, and tested the residue of these decanted and evaporated solvents (quinia, beberia, strychnia), and examined the aqueous solution of the

material, or one of the filtered alkaline liquids, in the usual way for acidulous radicals (citric, tartaric, sulphuric, hypophosphorous): Or he has modified his methods to include search for some "scale preparation" which his special knowledge tells him has been newly introduced to, or is rare in, pharmacy.

*In the case of liquids*, the solvents as well as the dissolved matters claim attention. A few drops are evaporated to dryness on platinum foil to ascertain if solid matter of any kind is present; the liquid is tested by red and blue litmus-paper to ascertain if free alkalies, free acids, or neither are present; a few drops are heated in a test-tube and the odor of any vapor noticed, a piece of glass tubing bent to a right angle being, if necessary, adapted to the test-tube by a cork, and some of the distilled liquid collected and examined; finally, the usual group-reagents for the several basylous and acidulous radicals are consecutively applied.

Proceeding in this way the student, who has already had some experience in pharmacy, will not be likely to overlook such solvents as water, acids, alkalies, alcohol, glycerine, ether, chloroform, benzole, fixed oils, and essential oils, or to miss the substances which these menstrua may hold in solution. He will probably also recognize such liquids as carbolic acid, formic acid, lactic acid, methylic alcohol, aldehyd, aniline, nitrobenzol. He must not, however, suppose that he will always be able to qualitatively analyze, say, a bottle of medicine; for the various infusions, decoctions, tinctures, wines, syrups, liniments, confections, extracts, pill-masses, and powders contain vegetable matters most of which at present are quite beyond the reach of the analyst. Neither the highest skill in analysis nor the largest amount of experience concerning the odor, appearance, taste, and uses of drugs is sufficient for the detection of all these vegetable matters. Skill and experience combined, however, will do much, and in most cases even so difficult a task as the one just mentioned be accomplished with reasonable success. Obviously, qualitative analysis alone will not enable the experimenter to produce a mixture of substances similar to that analyzed; to this end recourse must be had to quantitative analysis, a subject reserved for subsequent consideration.

Natural fluids, as "Milk" and "Urine" (*vide* Index), admit of special analytical treatment.

*Gas-analysis*, or *Eudiometry* (from εὐδία, *eudia*, calm air, and μέτρον, *metron*, a measure, in allusion to the *eudiometer*, an instrument used in measuring the proportion and, as the early chemists thought, the salubrity of the gases of the air), is a branch of experimental investigation, chiefly of a quanti-

tative character, concerning which information must be sought in other treatises. The analysis of atmospheric air from various localities, coal-gas, and gases obtained in chemical researches, involves operations which are scarcely within the sphere of Chemistry applied to Medicine. Beyond the recognition, therefore, of oxygen, hydrogen, nitrogen, carbonic, sulphurous, and hydrosulphuric acid gases, the experimental considerations of the chemistry of gaseous bodies may be omitted. Their study, however, should not be neglected, as existing conceptions of the constitution of chemical substances are largely dependent on the observed relations of the volumes of gaseous compounds to their elements. See previous paragraphs, pp. 36 to 43, 54 to 58, and 159. The best single work on this latter part of the subject is a small book by Hofmann, "Introduction to Modern Chemistry."

*Spectral Analysis.*—It may be as well to state here that the preliminary and final examinations of minute quantities of solid matter may, in certain cases, profitably include their exposure to a temperature at which they emit light, the flame being physically analyzed by a spectroscope. A spectroscope consists essentially of a prism to decompose a ray of light into its constituent colors, with tubes and lenses to collect and transmit the ray or rays to the eye of an observer. The material to be examined is placed on the end of a platinum wire, which is then brought within the edge of a spirit-lamp or other smokeless flame; volatilization, attended usually in the case of a compound by decomposition, at once occurs, and the whole flame is tinged with a characteristic hue. A flat ribbon of rays is next cut off by bringing near to the flame a brass tube, the cap of which is pierced by a narrow slit. At the other end of the tube, at focal distance for parallel rays, is a lens, through which the ribbon of light passes to a prism; the prism decomposes the ribbon, spreading out its constituent colors like a partially opened fan, and the colored beam or *spectrum* thus produced is then examined by help of a telescope attached by a movable joint to the stand which carries the prism and object-tube. Sodium compounds, under these circumstances, give yellow light only, indicated by a double band of light in a position corresponding to a portion of the yellow part of an ordinary solar spectrum. The potassium spectrum is mainly composed of a red and violet band; lithium a crimson, and, at very high temperatures, a blue band. Most of the other elements give equally characteristic spectra.

By aid of a combined microscope and spectroscope (microspectroscope) the color of colored fluids can be analyzed.

# CHEMICAL TOXICOLOGY.

In cases of criminal and accidental poisoning, the substances presented to the chemical analyst for examination are usually articles of food, medicines, vomited matters, or the liver, kidney, intestines, stomach and contents, removed in course of post-mortem examination. In these cases some special operations are necessary before the poison can be isolated in a state of sufficient purity for the application of the usual tests; for in most instances the large quantity of animal and vegetable, or, in one word, organic matter present, prevents or masks the characteristic reactions on which the tests are founded. These operations will now be described;* they form the chemical part of the subject of Toxicology (τοξικὸν, *toxicon*, poison, and λόγος, *logos*, discourse).

Substances occurring in the form of an apparently definite salt or unmixed with organic matter need no special treatment, they are analyzed by the ordinary methods already given, attention being restricted to poisonous compounds only.

---

## EXAMINATION OF AN ORGANIC MIXTURE SUSPECTED TO CONTAIN:—MERCURY, ARSENICUM, ANTIMONY, LEAD, COPPER, OR ZINC; SULPHURIC ACID, NITRIC ACID, HYDROCHLORIC ACID, OXALIC ACID, OR HYDROCYANIC ACID; CAUSTIC ALKALIES; PHOSPHORUS; STRYCHNIA, MORPHIA, OR OTHER POISONOUS ALKALOIDS.

### *Preliminary Examination.*

*Odor, Appearance, Taste.*—Smell the mixture with the view of ascertaining the presence or absence of any notable quantity of free hydrocyanic acid. Look carefully for any small solid particles, such as arsenic, corrosive sublimate, or verdigris, and for any appearance which may be regarded as abnormal, any character unusual to the coffee, tea, beer, medicine, vomit, coats of stomach, kidney, liver, or other organ, tissue, or solid matter under examination.

---

* Materials for these experiments are readily obtained for educational purposes by dissolving the poison in infusions of tea, coffee, porter, or in water to which some mucilage of starch, or linseed-meal, pieces of bread, potato, and fat have been added.

*Poisonous quantity of acid.*—Add to a small portion some solution of carbonate of sodium, with the view of ascertaining by strong effervescence the presence of any large, poisonous quantity of sulphuric, nitric, or hydrochloric acid (p. 486).

*Poisonous quantity of alkali.*—If so excessively alkaline as to require the addition of a very large quantity of acid before neutralization is effected, a noxious quantity of a corrosive or caustic alkali is present. Whether soda or potash is ascertained by the usual tests.

*Special instructions* may induce the operator to suspect the presence of one particular poison. Direct examination for the latter may then be made, either at once, if the substance has an aqueous character, or when filtration or treatment with warm hydrochloric or acetic acid has afforded a more or less colorless liquid.

*Fluids.*—A vomit or the contents of a stomach, if set aside in a long narrow vessel (test-glass or ale-glass), or, better, exposed on a filter during a night, will often yield a more or less clear limpid portion at the bottom or top of the solid matter. This fluid (separated by a pipette or otherwise) will sometimes respond to tests without further preparation, and always requires less preparatory treatment than a semisolid mixture. If none passes through a filter, a portion often collects in the upper part.

*General procedure.*—If the preliminary examination does not indicate the method to be pursued, proceed as follows, treating a portion (not more than one-fourth) of the mixture for the poisonous metals, another for the acids, and a third for alkaloids, reserving the remainder for any special experiments which may suggest themselves in the course of analysis.

### *Examination for Mercury, Arsenicum, Antimony, Lead, Copper, Zinc.*

If a liquid, acidulate with hydrochloric acid and boil for a short time. If solid or semisolid, cut up the matter into small pieces, add enough water to form a fluid mixture, stir in ten or twenty per cent. of ordinary liquid hydrochloric acid, and boil until, from partial aggregation and solution of the solid matter, filtration can be easily effected.

Heat a portion of the clear liquid with a thin piece of bright pure copper or copper gauze, about an inch long and a quarter of an inch broad, for about ten or twenty minutes; metallic *mercury*, *arsenicum*, or *antimony* will be deposited on the copper, darkening it considerably in color. Pour off the liquid

from the copper, carefully rinse the latter with a little cold water, dry the piece of metal by holding it over or near a flame (using fingers, not tongs, or it may become sufficiently hot for loss of mercury or arsenicum to occur by volatilization), introduce it into a narrow test-tube or piece of glass tubing closed at one end, and heat the bottom of the tube in a flame, holding it horizontally that the upper part of the tube may be kept cool, and partially closing the mouth with the finger to prevent escape of vapor. Under these circumstances any *Mercury* will volatilize from the copper and condense on the cool part of the tube in a ring or patch of white sublimate, readily aggregating into visible globules on being pressed by the side of a thin glass rod inserted into the tube; *Arsenicum* will volatilize from the copper, and, absorbing oxygen from the air in the tube, condense on the cool part of the glass in a ring or patch of white sublimate of arsenic (gray or even darker if much arsenicum as well as arsenic be present), not running into globules when rubbed, but occurring in small crystals, the characteristic octahedral form of which is readily seen by aid of a good hand lens, or the lower part of a microscope; *Antimony* volatilizes from the copper, if strongly heated, and, absorbing oxygen, immediately condenses as a slight white deposit close to the metal.

*Confirmatory Tests.* — 1. Nothing short of the production of globules should be accepted as evidence of the presence of mercury. It will usually have existed as corrosive sublimate.

2. To confirm indications of the presence of arsenicum, a portion of the acid liquid may be subjected to the hydrogen tests (pp. 173, 174); or the tube containing the white crystalline arsenic may be broken, and the part on which the sublimate occurs boiled for some time in water, and the hydrosulphuric-acid, ammonio-nitrate-of-silver, and ammonio-sulphate-of-copper tests (pp. 174, 177) applied to the aqueous solution.

3. For antimony, a portion of the acid liquid must always be introduced into the hydrogen apparatus with the usual precautions. (*Vide* p. 185.)

*Note.*—Before finally concluding that arsenicum is absent from a fluid, the latter should be warmed with a little sulphurous acid or oxalic acid, and ordinary tests then again applied; for arsenic acid and other arseniates are not readily affected by the usual reagents for arsenicum.

For *Lead* or *Copper*, pass hydrosulphuric acid gas through the clear acid liquid for some time, warming the liquid if no precipitate is produced, or diluting and partially neutralizing the acid by ammonia if much acid has been added. Collect on a filter any black precipitate that may have formed; wash, dissolve in a few drops of aqua regia, dilute, and apply the

tests, such as ammonia for copper, sulphuric acid for lead, and any other of the ordinary reagents (pp. 193, 214).

*Copper may often be at once detected* in a small quantity of acidulated liquid by immersing the point of a penknife or a piece of bright iron wire—a deposit of copper, in its characteristic color, quickly or slowly appearing, according to the amount present (p. 193).

*Zinc.*—To the acid liquid through which sulphuretted hydrogen has been passed, add excess of ammonia (or to the original acid fluid add excess of ammonia, and then sulphydrate of ammonium); a precipitate falls which may contain alumina, phosphates, and zinc; it is usually blackish from the presence of sulphide of iron. Collect the precipitate on a filter, wash, dissolve in a little hydrochloric acid, add a few drops of nitric acid, boil, pour in excess of ammonia, filter, and test the filtrate with sulphydrate of ammonia—a white precipitate indicates zinc.

### *Examination for Mineral Acids, Oxalic Acid, or Hydrocyanic Acid.*

To detect *Hydrochloric*, *Nitric*, or *Sulphuric Acid* in any liquid containing organic matter, dilute with water and apply to small portions the usual tests for each acid, disregarding indications of small quantities. (*Vide* pp. 269, 289, 311.)

Excessive sourness, copious evolution of carbonic acid gas on the addition of carbonate of sodium, and abundant evidence of acid on applying the various tests to small portions of the fluid presented for analysis, collectively form sufficient evidence of the occurrence of a poisonous amount of either of the three common mineral acids. Small quantities of the hydrochloric, nitric, and sulphuric radicals occurring as metallic salts or acids, are common normal constituents of food, hence the direction to disregard insignificant indications. If the fluid under examination be a vomit or the contents of a stomach, and an antidote has been administered, free acid will not be found, but, instead, a large amount of corresponding salt.

For *Oxalic Acid*, filter or strain a portion of the liquid, if not already clear, and add solution of acetate of lead so long as a precipitate occurs; collect the precipitate, which is partly oxalate of lead, on a filter, wash, transfer it to a test-tube or test-glass, add a little water, and pass hydrosulphuric gas through the mixture for a short time; the lead is thus converted into the insoluble form of sulphide, while oxalic acid is set free in the solution. Filter, boil to get rid of hydrosulphuric gas, and apply the usual tests for oxalic acid (see p. 317) to the clear filtrate.

The contents of a stomach containing oxalic acid are often of a dark-brown color with a tinge of green (altered blood and mucus), and the viscid mixture generally, though slowly, affords some clear, limpid, almost colorless liquid by filtration or standing.

For *Hydrocyanic Acid*, the three chief tests may be applied at once to the liquid or semi-liquid organic mixture, whether it has an odor of hydrocyanic acid or not. First: half fill a small porcelain crucible with the material, add eight or ten drops of strong sulphuric acid, stir gently with a glass rod, and invert over the mouth of the crucible a watch-glass moistened with a small drop of solution of nitrate of silver; a white film on the silver solution is probably cyanide of silver, formed by the action of the gaseous hydrocyanic acid on the nitrate of silver. Second: prepare a small quantity of the organic mixture as before, slightly moistening the centre of the watch-glass with solution of potash; here again the heat generated by the action of the strong acid is sufficient to volatilize some of the hydrocyanic acid, which, reacting on the potash, forms cyanide of potassium. On removing the watch-glass and stirring into it successively solution of a ferrous salt, a ferric salt, and hydrochloric acid, flocks of Prussian blue are produced if hydrocyanic acid is present. Third: proceed as before, moistening the watch-glass with sulphydrate of ammonium; after exposure to the hydrocyanic gas for five or ten minutes, add a drop of solution of ammonia, evaporate to dryness at a low temperature, and add a drop of hydrochloric acid and of solution of perchloride of iron; a blood-red color, due to sulphocyanate of iron, is produced if cyanogen is present.

If the above reactions are not well marked, the organic mixture may be carefully and slowly distilled in a small retort, the neck of which passes into a bottle and dips beneath the surface of a little water, at the bottom of the bottle, and the reagents then applied to separate portions of the distillate.

The examination of organic mixtures for hydrocyanic acid must be made without delay, as the poison soon begins to decompose, and in a day or two is usually destroyed.

### *Examination for Phosphorus.*

A paste containing phosphorus is commonly employed for destroying vermin. In cases of poisoning the phosphorus is commonly in sufficient quantity to be recognized by its characteristic unpleasant smell. A stomach in which it occurs not unfrequently exhibits slight luminosity if opened in a dark room. When the phosphorus is too small in quantity or too much diffused to afford this appearance, a portion of the material is placed in a flask, water acidulated by

sulphuric acid added, a long wide glass tube fitted to the neck of the flask by a cork, and the mixture gently boiled. If phosphorus is present (even 1 part in 2,000,000, according to De Vry) the top of the column of steam as it condenses in the tube will appear distinctly phosphorescent when viewed in a dark room. From its liability to oxidation phosphorus cannot be detected after much exposure of an organic mixture to air.

### *Examination for Strychnia and Morphia.*

*Strychnia.*—If solid or semisolid, digest the matter with water and about 10 per cent. of hydrochloric acid till fluid, filter, evaporate to dryness over a water-bath. If the organic mixture is already liquid, it is simply acidulated with hydrochloric acid and evaporated to dryness. The acid residue is next treated with spirit of wine as long as anything is dissolved, the filtered tincture evaporated to dryness over the water-bath, and the residue digested in water and filtered. This slightly acid aqueous solution must now be rendered alkaline by ammonia, and well shaken in a bottle or long tube with about half an ounce of chloroform, and set by till the chloroform has subsided. The chloroform (which contains the strychnia) is then removed by a pipette, the presence of any aqueous liquid being carefully avoided, and evaporated to dryness in a small basin over a water-bath, the residue moistened with concentrated sulphuric acid, and the basin kept over the water-bath for several hours. (It is highly important that the sulphuric acid used in this operation should be free from nitrous compounds. Test the acid, therefore, by adding powdered sulphate of iron, which becomes pink if nitrous bodies are present. If these are found, the acid should be purified by strongly heating with sulphate of ammonium, seventy or eighty grains to a pint.) The charred material is exhausted with water, filtered, excess of ammonia added, the filtrate shaken with about a quarter of an ounce of chloroform, the mixture set aside for the chloroform to separate, and the chloroform again removed. If, on evaporating a small portion of this chloroform solution to dryness, adding a drop of sulphuric acid to the residue, and warming, any darkening in color or charring takes place, the strychnia is not sufficiently pure for chemical detection; in that case the rest of the chloroform must be removed by evaporation, and the residue redigested in warm sulphuric acid for two or three hours. Dilution, neutralization of acid by ammonia, and agitation with chloroform are again practised, and the residue of a small portion of the chloroform solution once more tested with sulphuric acid. If charring still occurs, the treatment must be repeated

a third time. Finally a part of the chloroform solution is taken up by a pipette, and drop after drop evaporated on one spot of a porcelain crucible-lid until a fairly distinct dry residue is obtained. A drop of sulphuric acid is placed on the spot, another drop placed near, a minute fragment of red chromate of potassium placed in the second drop, and when the acid has become tinged with the chromate, one drop drawn across the other; the characteristic evanescent purple color is then seen, if strychnia is present. Other tests (*vide* p. 390) may be applied to similar spots.

This is Girdwood and Rogers's method for the detection of strychnia when mixed with organic matter. It is tedious but trustworthy, and, though apparently complicated, very simple in principle; thus: strychnia is soluble in acidulated water or alcohol, or in chloroform, readily removed from an alkaline liquid by agitation with chloroform, and not charred or otherwise attacked when heated to 212° F. with sulphuric acid: much of the organic matter of the food is insoluble in water: of that soluble in water, much is insoluble in alcohol; and of that soluble in both menstrua, all is charred and destroyed by warm sulphuric acid in a shorter or longer time.

*Morphia, and the Meconic Acid with which it is associated in Opium.*—To the liquid or the semifluid mixture warmed for some time in a small quantity of acetic acid, filtered, and concentrated if necessary, add solution of acetate of lead until no further precipitate is produced. Filter and examine the *precipitate* for meconic acid, reserving the *filtrate* for the detection of morphia.

*The Precipitate.*—Wash the *precipitate* (meconate of lead, etc.) with water, place it in a test-tube or test-glass with a small quantity of water, pass hydrosulphuric acid gas through the mixture for a short time, filter, slightly warm in a small basin, well stirring to promote removal of excess of the gas, and add a drop of neutral solution of perchloride of iron; a red color, due to the formation of meconate of iron, is produced if meconic acid is present. This color is not destroyed on boiling the liquid, as is the case with ferric acetate, nor is it bleached by solution of corrosive sublimate, thus distinguishing it from the ferric sulphocyanate. It is discharged by hydrochloric acid.

*The Filtrate.*—The solution from which meconic acid has been removed by acetate of lead is evaporated to a small bulk over a water-bath, excess of carbonate of potassium added, and evaporation continued to dryness. The residue is then treated with alcohol, which dissolves the morphia. The alcoholic solution evaporated similarly may leave the morphia sufficiently

pure for the application of the usual tests (*vide* page 382) to small portions of the residue. If no reaction is obtained, add a drop of sulphuric acid and a little water to the residue and shake with ether, in which the salt of morphia is insoluble. The treatment with ether may be repeated until nothing more is removed, the acid aqueous liquid saturated with carbonate of potassium, the mixture evaporated to dryness, the residue digested in alcohol, filtered, and portions of the alcoholic liquid evaporated to obtain spots of morphia for the application of the ordinary tests.

If much organic matter is believed to remain in the filtrate after the acetate of lead treatment, or if a considerable excess of acetate of lead has been employed, the filtered liquid should be subjected to a current of sulphuretted hydrogen until no more sulphide of lead is precipitated, the mixture filtered, and the filtrate, with the washings from the sulphide of lead, evaporated to a small bulk, excess of carbonate of potassium added, the whole well mixed and agitated with twice or thrice its bulk of a mixture of ether and acetic ether (ether alone might not dissolve the morphia). On standing the ethereal liquid rises to the surface; it is carefully removed, evaporated to dryness, and the residue tested or further purified in the manner described in the preceding paragraph.

The examination for morphia must be conducted with great care, and with as large a quantity of material as can be spared; for its isolation from other organic matter is an operation of considerable difficulty, especially when only a minute proportion of alkaloid is present. Fortunately the detection of meconic acid does not include similar difficulties; and, as its reactions are quite characteristic, its presence is held to be strong evidence of the existence of opium in an organic mixture.

### *Examination for other Poisonous Alkaloids.*

*Stas's Process.*—Minutely subdivide any solid matter; to this and the liquid portion of the vomit, etc., add about twice their weight of the strongest spirit of wine containing sufficient tartaric acid to fairly acidify the mixture. Digest the whole in a flask at a temperature of 150° or 160° F.; set aside to cool; filter. The solution, which will contain the whole of the alkaloid, should then be evaporated to dryness *in vacuo*, or at all events at a temperature not exceeding 100° F., lest volatile alkaloids should be dissipated. The residue is next exhausted with cold anhydrous alcohol; filtered; and the filtrate evaporated to dryness with the precautions already stated. The extract is dissolved in a very small quantity of water, treated with

The following is Brunner's Tabular Scheme for the separatio

TABLE OF REACTION

| Absorbed by Ether from Acid Solution.* | | | |
|---|---|---|---|
| With Tannin. | | | |
| Precipitable. | | Not precipitable. | |
| COLCHICINA. | DIGITALIN. | PICROTOXIN. | |
| Solution *yellow;* nitric acid colors it *violet.*<br><br>If the nitric acid solution is diluted and made alkaline with NaHO, an *orange-red* coloration is produced. | Mixed with solution of ox-gall and concentrated sulphuric acid a fine *red* zone is produced at first, afterwards a *red* liquid.<br>———<br>Dissolved in concentrated sulphuric acid and a trace of bromine added, a violet-red coloration appears. | The colorless solution in water containing soda is reduced by Fehling's copper solution. | |
| * Small quantity of Atropia. | | | |

| With co | |
|---|---|
| In the cold. | |
| Rose-red. | Brown |
| BRUCIA. | DELP |
| Dissolves in concentrated nitric acid with a *light-red* color, which on warming changes to *yellow:* if then chloride of tin is added, a *blue-violet* color appears. | With trated s acid a mine w *let-red.* same c appears porating phospho<br><br>*Aconi* solves *brown-r* in sulp acid. |

excess of powdered bicarbonate of sodium or potassium, and well shaken with five or six times its volume of pure ether (with perhaps a little acetic ether). This ethereal liquid contains the alkaloid. Small portions should be evaporated in watch-glasses and tasted, or tested physically and chemically, according as the knowledge of collateral circumstances by the operator, or his experience, or such reactions as are recorded on pp. 392–399, may suggest. (See Table.)

If a volatile alkaloid (conia, nicotia, hyoscyamia, lobelina) is indicated, the ethereal solution, which may still contain animal matter, is removed, agitated with aqueous solution of potash, decanted, and shaken with pure diluted sulphuric acid. On standing, the aqueous portion, containing the alkaloid as acid sulphate, subsides; the upper ethereal portion containing the animal matter is rejected; the acid aqueous liquid is made alkaline with caustic potash or soda; ether added; well shaken; the ethereal liquid decanted, evaporated to dryness *in vacuo*, or at a low temperature; and (to get rid of all traces of ammonia) again moistened with ether and dried. The residue is now tested for the suspected alkaloid by taste, smell, and the application of appropriate reagents (pp. 392–399). (See Table.)

If a non-volatile alkaloid (aconitia, atropia, brucia, colchicia, emetia, physostigmia. solania, veratria, as well as morphia, codeia, and strychnia) is indicated, further purification is effected by decanting ethereal liquid from the lower aqueous solution of bicarbonate of sodium, removing the ether by evaporation, digesting the residue in alcohol, filtering, evaporating the alcohol, treating the residue with dilute sulphuric acid, setting aside for a few hours, filtering, concentrating, adding powdered carbonate of potassium, and finally anhydrous alcohol. The alcoholic liquid, on evaporation, yields the alkaloid in a fit state for testing in the manner already stated. (See also annexed Table.)

*Sonnenschien's Process.*—Digest with diluted hydrochloric acid, evaporate to the consistence of syrup, dilute, set aside for some hours, filter. Add solution of phosphomolybdic acid so long as any precipitate falls or cloudiness occurs; collect the precipitate on a small filter; wash it with water containing phosphomolybdic and nitric acid, and, while still moist, place it in a flask. Decompose this compound of phosphomolybdic acid and alkaloid by adding caustic baryta until the stirred mixture is distinctly alkaline. Distil off volatile alkaloids, condensing and collecting by help of a long tube, so bent that the apparatus shall act as a retort, the end of the tube being attached to a

bulb or a series of bulbs containing dilute hydrochloric acid. The acid liquid evaporated gives a residue of hydrochlorates of alkaloids. The latter will afford characteristic reactions with the tests for the suspected alkaloid, and, on being moistened with baryta-water and warmed, will afford fumes of volatile alkaloids whose odor is usually characteristic. The residue in the flask will contain non-volatile alkaloids. It is treated with carbonic acid gas to neutralize and precipitate the excess of baryta as insoluble carbonate of barium; the mixture is evaporated to dryness over a water-bath; and the residue digested in alcohol. The alcoholic solution evaporated generally yields the alkaloids in a fit state for testing.

Phosphomolybdic acid forms with ammonia, in acid solutions, a remarkably insoluble compound, and it comports itself in a similar manner with those compounds which are analogous to ammonia—the nitrogenized organic bases—consequently forming an excellent reagent for their detection. It may be prepared in the following manner: Molybdate of ammonium is precipitated by phosphate of sodium; the yellow precipitate, having been washed, is diffused through water, and heated with sufficient carbonate of sodium to dissolve it. The solution is then evaporated to dryness, and calcined to drive off the ammonia. In case any of the molybdic compound be reduced by this operation, the residue must be moistened with nitric acid and again calcined. The dry mass is then dissolved in cold water, the solution strongly acidulated with nitric acid, and water added until ten parts of the solution contain one of the dry salt. The liquid, which is of a golden-yellow color, must be preserved from ammoniacal fumes. It precipitates all the alkaloids (with the exception of urea) when a mere trace only is present. The precipitates are yellow, generally flocculent, insoluble in water, alcohol, ether, and the dilute mineral acids, with the exception of phosphoric acid. Nitric, acetic, and oxalic acids, concentrated and boiling, dissolve them. These compounds are decomposed by the alkalies, certain metallic oxides, and the alkaline salts, which separate the alkaloid. To give an idea of the sensitiveness of this reagent, it may be stated that the 0.000071 gramme of strychnia gives an appreciable precipitate with one cubid centimetre of the solution of phosphomolybdic acid.

## ANTIDOTES.

*Vide* "Antidote" in the Index.

## QUESTIONS AND EXERCISES.

878. In examining food and similar matter for poison, why must not the ordinary tests for the poison be at once applied?

879. What preliminary operations should be performed on a vomit in a case of suspected poisoning?

880. How would you proceed in searching for corrosive sublimate in wine?

881. By what series of operations would you satisfy yourself of the presence or absence of arsenic in the contents of a stomach?

882. Describe the treatment to which decoction of coffee should be subjected in testing it for tartar-emetic.

883. State the method by which the occurrence of lead in water is demonstrated.

884. Give a process for the detection of copper in jam.

885. How would you detect zinc in a vomit?

886. How may the presence of a poisonous quantity of sulphuric acid in gin be proved?

887. In examining ale for free nitric acid what reactions would be selected?

888. Show how you would conclude that a dangerous quantity of hydrochloric acid had been added to cider.

889. Describe the manipulations necessary in testing for hydrocyanic acid in the contents of a stomach.

890. By what method is oxalic acid discovered in infusion of coffee?

891. How is the phosphorus detected in organic mixtures?

892. Give the process by which strychnia is isolated from partially digested food.

893. Mention the experiments by which the presence of laudanum in porter is demonstrated.

894. Name the appropriate antidotes in cases of poisoning by: *a*, alkaloids; *b*, antimonials; *c*, arsenic; *d*, barium salts; *e*, copper compounds; *f*, hydrochloric acid; *g*, hydrocyanic acid; *h*, preparations of lead; *i*, corrosive sublimate; *j*, nitric acid; *k*, oxalic acid; *l*, salts of silver; *m*, oil of vitriol; *n*, tin liquors; *o*, zinc solutions; *p*, carbolic acid.

---

# EXAMINATION OF MORBID URINE AND CALCULI.

The various products of the natural and continuous decay of animal tissue and the refuse matter of food are eliminated from the system chiefly as fæces, urine, and expired air. Air exhaled from the lungs carries off from the blood much carbon (about 8 ounces in 24 hours), in the form of carbonic acid gas, and some aqueous vapor—the latter, together with a small amount of oily matter, also escaping by the skin. Directing the breath to a cold surface renders moisture evident; and breathing through a tube into lime-water demonstrates the presence of a considerable quantity of carbonic acid gas. The fæces consist mainly of the insoluble *débris* of the system, the soluble matters and water forming the urine. These excretions vary con-

siderably, according to the food and general habits of the individual and external temperature. But in disease the variations become excessive; their detection by the medical practitioner, or by the pharmacist for the medical practitioner, is therefore a matter of importance.

A complete analysis of feces, urine, or expired air cannot be performed in the present state of our knowledge. Nor can even a partial analysis of feces or air be made with sufficient ease and rapidity to be practically available in medical diagnosis. But with regard to urine, certain abnormal substances and abnormal quantities of normal constituents may be chemically detected in the course of a few minutes by any one having already some knowledge of chemical manipulation.

*Healthy human urine* contains, in 1000 parts, 957 of water, 14 of urea, 1 of uric acid, 15 of other organic matter, and 13 of inorganic salts. The amount passed in twenty-four hours varies from two to three pints in an adult, and its specific gravity, if healthy, will range from 1.015 to 1.025. The acidity of urine Thudichum considers to be due to *cryptophanic acid*, $H_2C_5H_7NO_5$.

## Examination of Morbid Urine for Albumen, Sugar, Bile, and Excess of Urea; and Urinary Sediment for Urates (or Lithates), Phosphates, Oxalate of Calcium, and Uric Acid.

*Albumen.*—To detect albumen, acidulate a portion of the clear urine in a test-tube with a few drops of acid (to keep phosphates in solution—nitric is best, acetic not so good) and boil; flocks or coagula will separate if albumen be present.

This experiment should first be made on normal urine containing a drop or two of solution of white of egg. The coagulum is white if it is only albumen, greenish if bile pigment be present, and brownish-red if the urine contain blood. The influence of acids and alkalies on the precipitation of albumen is noticed on page 450.

The occurrence of albumen in the urine may be temporary and of but little importance; or it may indicate the existence of a serious affection, known as Bright's disease.

*Sugar.*—To a portion of the clear urine in a test-tube add five or ten drops of solution of sulphate of copper; pour in solution of potash or soda until the precipitate first formed is redissolved; slowly heat the solution to near the boiling-point; a yellow, yellowish-red, or red precipitate (cuprous oxide) is formed if sugar be present.

This experiment should first be made on urine containing a drop or two of solution of grape-sugar (page 409). The hydrate of copper precipitated by the alkali is insoluble in excess of pure potash or soda, but readily dissolves if organic matter, especially sugar, be present. The copper salt may not contain iron.

Other tests may be applied if necessary (*vide* pages 409 *et seq.*).

A minute amount of sugar is said to occur in normal urine, and a

distinct trace is occasionally present. In larger quantities (often 5 per cent.) it is a characteristic constituent of the urine of diabetic patients, greatly increasing the specific gravity of the excretion. Small hydrometers (termed urinometers) are commonly employed for quickly and readily ascertaining the specific gravity of urine; they range from 1.000 to 1.050, the interval of 1.015 to 1.025 being marked as "H. S." or healthy state. (*Vide* "Specific Gravity" and "Hydrometers" in Index.)

*Bile.*—This is best detected by the general test (Pettenkofer's) described on page 457. Or a little of the urine may be placed on a white plate, and strong nitric acid dropped on it; a peculiar play of colors—green, yellow, violet, etc.—occurs if (the coloring matter of) bile be present.

*Excess of Urea.*—About one-third of the solid matter in the urine is urea. Its proportion varies considerably; but $1\frac{1}{2}$ per cent. may be regarded as an average amount. Concentrate urine slightly by evaporation in a small dish, pour the liquid into the test-tube, set the tube aside till cold, or cool it by letting cold water run over the outside, add an equal bulk of strong nitric acid and again set aside; scaly crystals of nitrate of urea are deposited more or less quickly.

With regard to the amount of urea in urine, it is impossible to sharply define excess or deficiency. If nitric acid gives crystals without concentration, excess is certainly present. A rough estimate may be formed by mixing a few drops of the urine and acid on a piece of glass and setting aside; the time which elapses before crystals form is an indication of the quantity in the specimen. The time will vary according to the temperature and state of moisture of the atmosphere; but with care some useful comparative results may in this way be obtained.

For methods of estimating the amount of urea in urine, and for drawings of the necessary apparatus, *vide* "Journal of the Chemical Society," 1874, p. 749; and 1877, i. pp. 534 and 538; also "Chemical News," Jan. 22, 1875.

*Tests.*—Urea in solution in water may be detected by the above reaction with nitric acid, and by the readiness with which it yields ammonia on being boiled with alkalies. In putrid urine its conversion into an ammoniacal salt has already been effected.

$$\underset{\text{Urea.}}{CH_4N_2O} + \underset{\text{Water.}}{2H_2O} = \underset{\text{Carbonate of ammonium.}}{(NH_4)_2CO_3}.$$

*Formula of Urea.*—The empirical formula of urea is $CH_4N_2O$. Its rational formula may be thus written:— $\left.\begin{matrix}(CO)'' \\ H_2 \\ H_2\end{matrix}\right\} N_2$; that is, it may be regarded as one of the organic bases already referred to, a primary diamine, in which the bivalent radical CO occupies the place of $H_2$. The other atoms of hydrogen may be displaced by various radicals, and many *compound ureas* be thus obtained.

*Artificial Urea.*—Urea may be prepared artificially by Willams' modification of Wöhler's method. Cyanide of potassium of the best commercial quality (containing about 90 per cent. of real cyanide), is fused at a very low red heat in a shallow iron vessel; red lead is added in small quantities at a time, the temperature being kept down by constant stirring. When the red lead ceases to cause further action, the mixture (cyanate of potassium and lead) is allowed to cool, the product finely powdered, exhausted with cold water, nitrate of barium added till no more precipitate (carbonate of barium) falls, the mixture filtered, and the filtrate treated with nitrate of lead so long as cyanate of lead is thrown down. The latter is thoroughly washed, and dried at a low temperature. Equivalent quantities of cyanate of lead and sulphate of ammonium, digested in a small quantity of water at a gentle heat and filtered, yield a solution from which urea crystallizes on cooling.

*Another Process.*—Basaroff has found that urea is produced when ordinary carbonate of ammonium is heated in hermetically sealed tubes to about 275° F. for a few fours. The same chemist had previously obtained urea by similarly heating pure carbonate of ammonium, so that the source of the urea in the former case is probably the carbonate of ammonium believed to occur in the carbonate (see page 92).

$$NH_4NH_2CO_2 - H_2O = CH_4N_2O.$$

## Urinary Sediments.

**Warm the sediment with the supernatant urine and filter.**

<table>
<tr><td colspan="3">Insoluble.<br><br>Phosphates, oxalate of calcium, and uric acid.<br>Warm with acetic acid, and filter.</td><td>Soluble.<br><br>Urates—of ammonium, calcium. or sodium, chiefly the latter.<br><br>They are redeposited as the liquid cools, and if sufficient in quantity may be further examined for ammonium, calcium, sodium, and the uric radical by the appropriate tests.</td></tr>
<tr><td colspan="2">Insoluble.<br><br>Oxalate of calcium and uric acid.<br>Warm with hydrochloric acid, filter.</td><td rowspan="2">Soluble.<br><br>Phosphates.<br>Add ammonia, white ppt. = phosphate of calcium, or ammonio-magnesium phosphate, or both.</td></tr>
<tr><td>Insoluble.<br><br>Uric acid.<br>Apply murexid test (p. 359).</td><td>Soluble.<br><br>Oxalate of calcium.<br>May be precipitated by ammonia.</td></tr>
</table>

*Notes.*—Urinary deposits are seldom of a complex character; the action of heat and acetic and hydrochloric acids generally at once indicates the character of the deposit, rendering filtration and precipitation unnecessary.

*The urates* are often of a pink or red color, owing to the presence of a pigment termed *purpurine;* hence the common name of *red gravel* for such deposits. *Purpurine* is soluble in alcohol, and may be removed by digesting a red deposit in that solvent. It is seldom necessary to determine whether the urate be that of ammonium, calcium, or sodium (see also Uric Acid, p. 359).

The *phosphate of calcium* and the *ammonio-magnesium phosphate* are usually both present in a phosphatic deposit, the magnesium salt forming the larger proportion. They may, if necessary, and if sufficient in quantity, be separated by collecting on a filter, washing, and boiling with solution of carbonate of sodium. The carbonates of calcium and magnesium thus formed are collected on a filter, washed, and dissolved in a drop or two of hydrochloric acid; chloride of ammonium, ammonia, and carbonate of ammonium are added, and the mixture boiled and filtered; any calcium originally present will then remain insoluble as carbonate of calcium, while any magnesium will be precipitated from the filtrate as ammonio-magnesium phosphate on the addition of phosphate of sodium, the mixture being also well stirred.——The chief portion of excreted phosphates is carried off by the feces, that remaining in the urine being kept in solution by the influence of acid phosphate of sodium, and frequently lactic acid.——Occasionally, an hour or two after a hearty meal, the urine becomes sufficiently alkaline for the phosphates to be deposited, and the urine when passed is turbid from their presence.——The ammoniacal constituent of the magnesium salt does not occur normally, but is produced from urea as soon as urine becomes alkaline.

*Oxalate of calcium* is seldom met with in excessive amounts, but very often in small quantities mixed with phosphates.

In one case of oxaluria the whole urine excreted by a patient in twenty-four hours furnished to the author only two-thirds of a grain of oxalate of calcium.

*Free uric acid* is in most cases distinctly crystalline and nearly always of a yellow, red, or brown color.

*Artificial Sediments.*—For educational practice, artificial deposits may be obtained as follows: 1. Rub up in a mortar a few grains of serpent's excrement (chiefly urate of ammonium) with an ounce or two of urine; this represents a sediment of urates. 2. Add a few drops of solution of chloride of calcium and of phosphate of sodium to urine; the deposit may be regarded as one of phosphates. 3. To an ounce or two of urine add very small quantities of chloride of calcium and oxalate of ammonium; the precipitate is oxalate of calcium. 4. To urine acidulated by hydrochloric acid add a little serpent's excrement; the sediment is uric acid.

*Other deposits* than the foregoing are occasionally observed. Thus *hippuric acid* ($HC_9H_8NO_3$), a normal constituent of human urine, and largely contained in the urine of herbivorous animals, is some-

times found associated with uric acid in urinary sediment, especially in that of patients whose medicine contains benzoic acid (p. 335). Its appearance, as observed by the aid of the microscope, is characteristic —namely, slender, four-sided prisms, having pointed ends. *Cystin* ($C_3H_7HSO_2$) (from κύστις, *kustis*, a bladder, in allusion to its origin) rarely occurs as a deposit in urine. It is not soluble in warm urine or dilute acetic acid, and scarcely in dilute hydrochloric acid, hence would be met with in testing for free uric acid. It is very soluble in ammonia, recrystallizing from a drop of the solution placed on a piece of glass in characteristic microscopic six-sided plates. *Organized sediments* may be due to the corpuscles of pus, mucus, or blood, fat globules, spermatozoa, cylindrical casts of the tubes of the kidneys, epithelial cells from the walls of the bladder, or foreign matters, such as fibres of wood, cotton, small feathers, dust; these are best recognized by the microscope, as will be seen by the following paragraphs and figures on the microscopic appearances of both crystalline and organized urinary sediments.

## Microscopic Examination of Urinary Sediments.

Urine containing insoluble matter is usually more or less opaque. For microscopical examination a few ounces should be set aside in a conical test-glass for an hour or two, the clear supernatant urine poured off from the sediment as far as possible, a small drop of the residue placed on a slip of glass and covered with a piece of thin glass and examined under the microscope with different magnifying-powers.

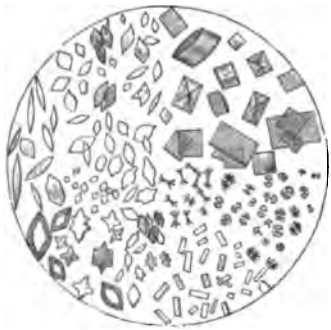
The respective appearances of the various crystalline and organized matters are given in the following figures, which were kindly drawn by H. B. Brady, F.R.S., from natural specimens (as seen with a two-third inch objective and No. 1 eye-piece, *i. e.*, magnified 60 diameters) in the collections of St. Bartholomew's Hospital, Dr. Sedgwick, W. W. Stoddart, F.C.S., Mr. Waddington, and the Author.

*Uric Acid* occurs in many forms, most of which are given in the first two figures. Flat, more or less oval crystals, sometimes attached to each other, their outline then resembling an 8, a cross, or a star, are common. Single and grouped quadratic prisms, aigrettes, spicula, and crystals recalling dumb-bells are met with. From urine acidulated by hydrochloric acid, square crystals, two opposite sides smooth and two jagged, are generally deposited; acidulated by acetic acid more typical forms are obtained. A drop of solution of potash or soda placed on a glass slip will dissolve a deposit of uric acid, a drop of any acid reprecipitating it in minute but characteristic crystals.

*Cystin* is very rarely met with as a urinary deposit; that from which the figure was taken was found in the urine of a patient in St. Bartholomew's Hospital. Lamellæ of cystin always assumes an hexagonal character; but the angles are sometimes ill defined and the plates superposed; in the latter case, a drop of solution of ammonia

placed on the glass at once dissolves the deposit, well-marked six-sided crystals appearing as the drop dries up.

Fig. 50.



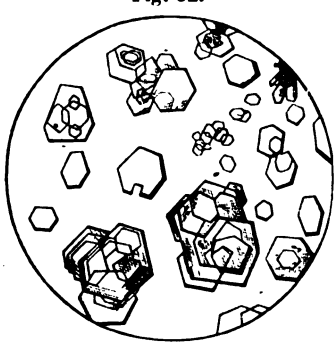
Uric Acid.

Fig. 51.

Uric Acid.

*Triple Phosphate* (phosphate of magnesium and ammonium) is deposited as soon as urine becomes alkaline, the ammoniacal constituent being furnished by the decomposition of urea. It occurs in

Fig. 52.



Cystin.

Fig. 53.

Triple Phosphate.

large prismatic crystals, forming a beautiful object when viewed by polarized light—sometimes also in ragged stellate or arborescent crystals, resembling those of snow. Both forms may be artificially prepared by adding a small lump of carbonate of ammonium to a few ounces of urine and setting aside in a test-glass.

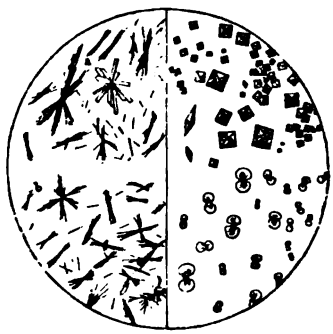
*Amorphous deposits* are either *earthy phosphates* (a mixture of phosphates of magnesium and calcium) or *urates* (of calcium, magnesium, ammonium, potassium, or sodium—chiefly the latter). They may be distinguished by the action of a drop of acetic acid placed near the sediment on the glass slip, the effect being watched under

the microscope; phosphates dissolve, while urates gradually assume characteristic forms of uric acid. Urates redissolve when warmed with the supernatant urine.

*Urates of Sodium and Magnesium*, though generally amorphous, occasionally take a crystalline form—bundles or tufts of small needles—as shown in the cut.

*Oxalate of Calcium* commonly occurs in octahedra requiring high magnifying power for their detection. The crystals are easily overlooked if other matters are present, but are more distinctly seen after phosphates have been removed by acetic acid. In certain aspects the smaller crystals look like square plates traversed by a cross. A dumb-bell form of this deposit is also sometimes seen, resembling certain forms of uric acid and the coalescing spherules of a much rarer sediment — carbonate of calcium. Oxalate of calcium is insoluble in acetic but soluble in hydrochloric acid. The octahedra are frequently met with in the urine of persons who have partaken of garden rhubarb; the crystals may often be deposited artificially (according to Waddington) by dropping a fragment of oxalic acid into several ounces of urine and setting aside for several hours.

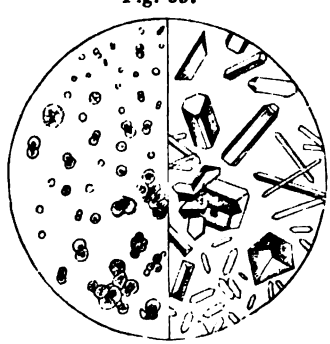
Fig. 54.



Urates, a, of Sodium, b, of Magnesium. Oxalate of Calcium.

*Carbonate of Calcium* is rarely found in the urine of man, but frequently in that of the horse and other herbivorous animals. Human urine containing carbonate of calcium often reddens litmus paper; and it is only after the removal, on standing, of the excess of carbonic acid that the salt is deposited. It consists of minute spherules, varying in size, the smaller ones often in process of coalescence. The dumb-bell form thus produced is easily distinguished from similar groups of uric acid or oxalate of calcium by showing a black cross in each spherule when viewed by polarized light. Acetic acid dissolves carbonate of calcium, liberating carbonic acid gas, with visible effervescence (under the microscope) if the slide has been previously warmed and a group of crystals be attacked.

Fig. 55.



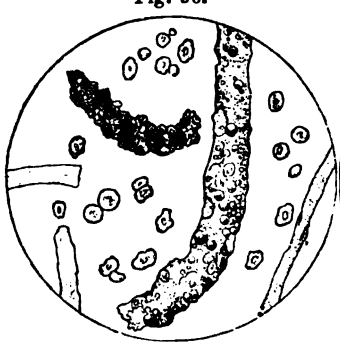
Carbonate of Calcium. Hippuric Acid.

*Hippuric Acid.*—The pointed rhombic prisms and acicular crystals are characteristic, and easily recognized. The broader crystals may possibly be mistaken for triple phosphate, and the narrower for certain forms of uric acid; but insolubility in acetic acid distinguishes them from the former, and solubility in alcohol from the latter. These tests may be applied while the deposit is under microscopic observation. An alcoholic solution of hippuric acid evaporated to dryness, and the residue treated with water, gives a solution from which characteristic crystalline forms of hippuric acid may be obtained on allowing a drop to dry upon a slip of glass.

The organized deposits in urine entail greater care in their determination, and usually require a higher magnifying power for their proper examination, than those of crystalline form. The figures are drawn to 230 diameters. The following notes will assist the observer.

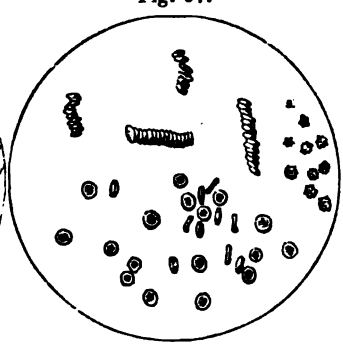
*Casts of uriniferous tubuli* are fibrinous masses of various forms, and often of considerable length—sometimes delicate and transparent, occasionally granular, and often beset with fat-globules. Epithelial *débris* are frequently present in urine in the form of nucleated cells, regular and oval when full, but angular and unsymmetrical when partially emptied of their contents—sometimes perfect, but more frequently a good deal broken up.

Fig. 56.



Epithelial Cells and Tubuli.

Fig. 57.



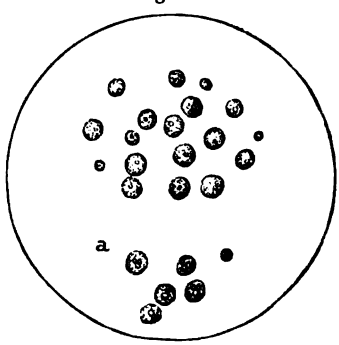
Blood Corpuscles.

*Blood* is easily recognized. Urine containing it is high-colored, and the corpuscles appear under the microscope as reddish circular disks, either single or laid together in strings resembling piles of coin. Their color and sometimes smaller size serve to distinguish them from pus-corpuscles. In doubtful cases a minute drop of blood taken from a finger by help of a needle should be diluted with water and used for comparison. After urine containing blood has stood for some time, the corpuscles lose their regular outline and become angular. (See *a* in the figure.) Day, of Geelong, tests for blood in urine, or in stains on clothing, by adding a few drops of a recently prepared alcoholic

solution of the inner unoxidized portions of guaiacum resin and then a small quantity of Robbin's aqueous or ethereal solution of peroxide of hydrogen, when a blue color results. "If the stain is on a dark-colored fabric, the moistened parts may be pressed with white blotting paper, when blue impressions will be obtained. Contact with many substances causes the blue reaction or oxidation of guaiacum; the peculiarity of blood is that it does not produce this effect unless peroxide of hydrogen or a similar 'antozonic' liquid is present. Bodies such as permanganate of potassium, whose oxygen is, apparently, in the form of ozone, also give rise to a blue color with guaiacum; peroxide of hydrogen and other compounds whose oxygen is in the opposite, positive, or, according to Schönbein, antagonistic condition, produce no such effect. It would seem as if blood or some other constituent of blood has the power of converting positive into negative oxygen, and thus bring about an effect which negative oxygen alone is able to produce; for of all substances which, like blood, do not alone cause guaiacum to become blue, blood is the only one that so affects 'antozonides' (themselves inactive) as to enable them to act as ozonides, that is to oxidize the guaiacum. Both the venous and arterial fluid from any red-blooded animal will produce this blue reaction. Fruit stains are darkened by ammonia, which does not alter the color of blood. Iron stains or iron-mould yield no color to water, whereas the red coloring-matter of blood is soluble in water. The peroxide of hydrogen should be free from more than a trace of acid."

The blood-corpuscles of ordinary animals are much smaller than those of man, but a $\frac{1}{16}$ or $\frac{1}{25}$ of an inch lens is necessary for proper differentiation (J. G. Richardson).

Fig. 58.

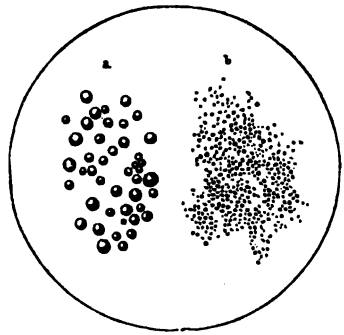



Pus Corpuscles.

*Pus and Mucus.*—Purulent urine deposits, on standing, a light-colored layer, easily diffused through the liquid by shaking. Acetic acid does not dissolve the sediment; and solution of potash, of official strength, converts it into a gelatinous mass. Under the microscope, pus-corpuscles appear rounded and colorless, rather larger than blood-disks, and somewhat granular on the surface. They generally show minute nuclei, which are more distinctly seen after treatment with acetic acid. (See the portion of the figure marked *a*.) Mucus possesses no definite microscopic characters, but commonly has imbedded in it pus, epithelium, and air-bubbles. Mucus is coagulated in a peculiar and characteristic manner by acetic acid: and this reaction, together with the ropy appearance it imparts to urine, prevents its being confounded with pus. Day's test for pus consists in adding a drop or two of oxidized

tincture of guaiacum to the urine or other liquid, when a clear blue color is produced. It is necessary to moisten dry pus with water before applying the test. The test liquid is made by exposing a saturated alcoholic solution of guaiacum to the air until it has absorbed a sufficient quantity of oxygen to give it the property of turning green when placed in contact with iodide of potassium. Day's test for mucus consists in the application, first, of oxidized tincture of guaiacum, which by itself undergoes no change in the presence of mucus, and then in the addition of carbolic acid or creasote, which quickly changes the color of the guaiacum to a bright blue. Neither carbolic acid nor creasote alone will render guaiacum blue. In testing for mucus on cloths, or when it is mixed with blood, it is necessary to use the carbolic acid pure, but when the mucus is in a liquid state it is better to use carbolic acid diluted with alcohol.

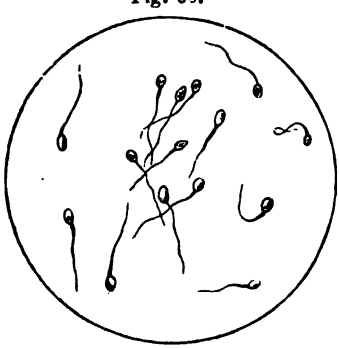
Fig. 59.




Fat Globules.

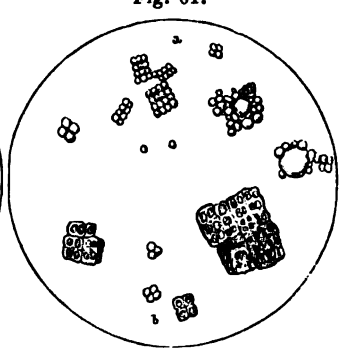
*Saliva.*—Saliva is an aqueous fluid containing less than 1 per cent. of solid matter, of which one-third is an albuminoid substance termed *ptyalin* (from πτύελον, *spittle*), a body that has power of converting starch into dextrin and grape sugar. Alkaline salts, including a trace of sulphocyanate of potassium, and calcareous compounds, are also present.

Fig. 60.



Spermatozoa.

Fig. 61.




Sarcina Ventriculi.

Day's test for saliva in urine, etc., is similar to that for mucus, with the exception that the blue reaction produced by the oxidized

tincture of guaiacum and alcoholic solution of carbolic acid is highly intensified by the addition of Robbin's aqueous or ethereal solution of peroxide of hydrogen.

*Fatty matter* occurs either as minute globules partially diffused through the urine (as shown at *a*, Fig. 59) or in more intimate emulsion (as at *b*). When present in larger quantity, it collects as a sort of scum on the surface after standing.

*Spermatozoa* are liable to escape notice, on account of their small size and extreme transparency. Suspected urine should be allowed to settle some hours in a conical test-glass, and the drop at the bottom examined under a high power. The drawing, Fig. 60, shows their tadpole-like appearance.

*Sarcina ventriculi* is an alga of a very rare occurrence in urine, though not unfrequent in the matters vomited during certain diseases of the stomach. The upper figures (*a*, Fig. 61) are copied from Dr. Thudichum's drawing (from urine); the larger fronds (*b*) are from vomited matter.

*Extraneous bodies*, such as hair, wool, or fragments of feathers, are often found in urinary deposits; and ludicrous mistakes have been made by observers not on their guard in respect to such casual admixtures.

## Examination of Urinary Calculi.

The term *calculus* is the diminutive of *calx*, a lime- or chalk-stone.

Knowledge of the composition of a calculus or urinary deposit affords valuable diagnostic aid to the physician; hence the importance of a correct analysis of these substances.

*Nature of Calculi.*—Urinary calculi have the same composition as unorganized urinary sediments. They consist, in short, of sediments that have been deposited slowly within the bladder, particle on particle, layer on layer, the several substances becoming so compact as to be less easily acted on by reagents than when deposited after the urine has been passed—the urates less readily soluble in warm water, the calcic phosphate insoluble in acetic acid until it has been dissolved in hydrochloric acid and reprecipitated by an alkali.

*Preliminary Treatment.*—If the calculus is whole, saw it in two through the centre, and notice whether it is built up of distinct layers or apparently consists of one substance. If the latter, use about a grain of the sawdust for the analysis; if the former, carefully scrape off portions of each layer, and examine them separately. If the calculus is in fragments, select fair specimens of about half a grain or a grain each, and reduce to a fine powder by placing on a hard surface and crushing under the blade of a knife.

*Analysis.*—Commence the analysis by heating a portion about the size of a pin's head, on platinum foil, in order to ascertain whether organic matter, inorganic matter, or both, are present. If both, the ash is examined for inorganic sub-

stances, and a fresh portion of the calculus for uric acid by the murexid test (In the absence of uric acid any slight charring may be considered to be due to indefinite animal matter.) If composed of organic matter only, the calculus will in nearly all cases be uric acid, the indications being confirmed by applying the murexid test in a watch-glass to another fragment, half the size of a small pin's head. If inorganic only, the ash on the platinum foil may be examined for phosphates, and a separate portion of the calculus for oxalates. Even a single drop of liquid obtained in any of these experiments may be filtered by placing it on a filter not larger than a sixpence and previously moistened with water, and adding three or four drops of water one after the other as each passes through the paper. If the calculus is suspected to contain more than one substance, boil about half a grain of the powder in half a test-tubeful of distilled water for a few minutes and pour it on a small filter; then proceed according to the following Table—

<table>
<tr><td colspan="2">Insoluble.<br><br>Phosphates, oxalate of calcium, and free uric acid.<br>Boil with two or three drops of hydrochloric acid, and filter.</td><td>Soluble.<br><br>Urates.<br>These will probably be redeposited as the solution cools. Small quantities may be detected by evaporating the solution to dryness. They are tested for ammonium, sodium, calcium, and the uric radical by the appropriate reagents.</td></tr>
<tr><td>Insoluble.<br><br>Uric acid. Apply the murexid test (p. 359).</td><td colspan="2">Soluble.<br><br>Phosphates and oxalate of calcium.<br>Add excess of ammonia, and then excess of acetic acid; filter.</td></tr>
<tr><td rowspan="2"></td><td>Insoluble.<br><br>Oxalate of calcium.</td><td>Soluble.<br><br>Phosphates.<br>They may be reprecipitated by ammonia.</td></tr>
</table>

*Varieties of Calculi.*—Calculi composed entirely of *uric acid* are common; a minute portion heated on platinum foil chars, burns, and leaves scarcely a trace of ash. The phosphates frequently occur together, forming what is known as the *fusible calculus*, from the readiness with which a fragment aggregates, and even fuses to a bead, when heated on a loop of platinum wire in the blowpipe-flame.

The phosphates may, if necessary, be further examined by the method described in connection with urinary deposits. Oxalate of calcium often occurs alone, forming a dark-colored calculus having a very rough surface, hence termed the *mulberry calculus*. Smaller calculi of the same substance are called, from their appearance, *hempseed calculi*. Calculi of *cystin* are rarely met with. *Xanthin* (from ξανθὸς, *xanthos*, yellow, in allusion to the color it yields with nitric acid) still less often occurs as a calculus. The earthy concretions, or *chalk-stones*, which frequently form in the joints of gouty persons, are composed chiefly of urates, the sodium salt being that most commonly met with. *Gall-stones*, or *biliary calculi*, occasionally form in the gall-bladder; they contain *cholesterin* (from χολή, *cholē*, bile, and στερεὸς, *stereos*, solid), a fatty substance of alcoholoid constitution, soluble in rectified spirit or ether, and crystallizing from such solutions in well-defined, square, scaly crystals. Phosphatic and other calculi of many pounds weight are occasionally found in the stomach and larger intestines of animals.

## QUESTIONS AND EXERCISES.

895. In breathing, how much carbon (in the form of carbonic acid gas) is exhaled from the lungs every 24 hours?

896. How may the presence of carbonic gas in expired air be demonstrated?

897. Mention an experiment showing the escape of moisture from the lungs during breathing.

898. State the method of testing for albumen in urine.

899. Give the tests for sugar in urine.

900. What is the average composition of healthy urine?

901. Give the tests for urea.

902. Write the rational formulæ of some compound ureas in which methyl or ethyl displaces hydrogen.

903. Describe an artificial process for the production of urea, giving equations.

904. Sketch out a plan for the chemical examination of urinary sediments.

905. A deposit is insoluble in the supernatent urine or in acetic acid; of what substance may it consist?

906. Which compounds are indicated when a deposit redissolves on warming it with the supernatent urine?

907. Name the salts insoluble in warmed urine but dissolved on the addition of acetic acid.

908. Mention the chemical characters of cystin. At what stage of analysis would it be recognized?

909. Describe the Microscopical appearance of the following urinary deposits:—

| | |
|---|---|
| Uric acid. | Tube-casts. |
| Cystin. | Epithelial *débris*. |
| Triple phosphate. | Blood. |
| Earthy phosphates. | Pus. |
| Urates. | Mucus. |
| Oxalate of Calcium. | Fat. |
| Carbonate of Calcium. | Spermatozoa. |
| Hippuric Acid. | Sarcina. |

Extraneous Bodies.

910. How are Day's tests for blood, pus, and saliva applied?

911. What is the general, physical, and chemical nature of urinary calculi?

912. How are urinary calculi prepared for chemical examination?

913. Draw out a chart for the chemical examination of urinary calculi.

914. Why is the "fusible calculus" so called, and what is its composition?

915. State the characters of "mulberry" and "hempseed" calculi.

916. What are the "chalk-stones" of gout, and "gall-stones" or "biliary calculi"?

---

# THE GALENICAL PREPARATIONS OF THE PHARMACOPŒIAS.

The preparation of Confections, Decoctions, Enemas, Extracts, Glycerines, Infusions, Inhalations, Juices, Liniments, Lozenges, Mixtures, Ointments, Pills, Plasters, Poultices, Powders, Spirits, Suppositories, Syrups, Tinctures, and Wines, includes a number of mechanical rather than chemical operations, and belongs to the domain of pure Pharmacy. The medical or pharmaceutical pupil will have had ample opportunity of practically studying those compounds before working at experimental chemistry, and will probably have prepared many of them according to the directions of the Pharmacopœias; if not, he is referred to the pages of the last edition of those works for details.

Among the extracts of the British Pharmacopœia, however, there are five (namely, those of Aconite, Belladonna, Hemlock, Henbane, and Lettuce) which are not simply evaporated infusions, decoctions, or tinctures, like most others, but are evaporated juices from which vegetable albumen, the supposed source of fermentation and decay, has been removed, and chlorophyll (the green coloring-matter of plant-juice) retained practically

unimpaired in tint. For educational practice either of the above-named five raw materials may be employed; but in order that attention may be concentrated on the process by which the extracts are prepared, rather than on any one of the extracts themselves, it suffices to make an extract of some ordinary green vegetable, such as cabbage or turnip tops. Bruise the green leaves of a good-sized cabbage in a mortar, and press out the juice; heat it gradually to 130° F., and remove the green flocks of chlorophyll which separate, by filtration through calico. When the liquor has all passed through the filter, set the chlorophyll aside for a time, heat the strained liquor to 200° F. to coagulate albumen; remove the latter by filtration and throw it away; evaporate the filtrate by a water-bath to the consistence of thin syrup; then add to it the chlorophyll, and, stirring the whole together assiduously, continue the evaporation at a temperature not exceeding 140° F., until the extract is of a suitable consistence for forming pills. A higher temperature than that indicated would cause the alteration of the chlorophyll to a dark-brown substance, any such extract used in pharmacy no longer having the green tint which custom and the British Pharmacopœia demand.

## QUESTIONS AND EXERCISES.

917. Enumerate the different classes into which official galenical preparations may be divided.

918. Describe the general process for the preparation of green extracts:—

Aconite. Hemlock.
Belladonna. Henbane.
Lettuce.

919. Why is vegetable albumen excluded in the preparation of green extracts?

920. How may chlorophyll be removed from vegetable juices, and again be introduced into their evaporated residues, without destroying its color?

921. For what reason is exposure of chlorophyll to a boiling temperature avoided in the manufacture of green extracts?

# THE CHEMICAL PREPARATIONS OF THE PHARMACOPŒIAS.

The process by which every official chemical substance is prepared has already been described, and the strict chemical character of the processes illustrated by experiments and explained by aid of equations. Should the reader, in addition, desire an intimate acquaintance with those details of manipulation on which the successful and economic manufacture of chemical substances depends, he is advised to prepare, if he has not done so already, a few ounces of each of the salts mentioned in the Pharmacopœias or commonly used in Pharmacy. An additional guide in these operations will be the Pharmacopœia itself.

The production of many chemical and galenical substances on a commercial scale can only be successfully carried on in manufacturing-laboratories and with some knowledge of the circumstances of supply and demand, value of raw material, and of by-products, etc.; for the technical preparation of such substances requires much knowledge beyond even a thorough acquaintance with chemistry. Still in the present day, commercial Chemistry and Pharmacy can best hope for success when founded on the working out of abstract scientific principles. The problem of manufacturing-success is now only solved with certainty by sound and wisely-applied science.

---

*Memorandum.* — The next subject of experimental study will be determined by the nature of the student's future pursuits. In most cases the operations of quantitative analysis will engage attention. These should be of a volumetric and gravimetric character; for details concerning them see the following pages.

# QUANTITATIVE ANALYSIS.

## INTRODUCTORY REMARKS.

*General Principles.*—The proportions in which chemical substances unite with each other in forming compounds are definite and invariable (p. 47). Quantitative analysis is based on this law. When, for example, aqueous solutions of a salt of silver and a chloride are mixed, a white curdy precipitate is produced containing chlorine and silver in atomic propertions, that is, 35.5 parts of chlorine to 108 of silver. No matter what the chloride or what the salt of silver, the resulting chloride of silver is invariable in composition. The formula AgCl is a convenient picture of this compound in these proportions. The weight of a definite compound being given, therefore, the proportional amounts of its constituents can be ascertained by simple calculation. Suppose, for instance, 8.53 parts of chloride of silver have been obtained in some analytical operation, this amount will contain 2.11 parts of chlorine and 6.42 of silver: for if 143.5 (the molecular weight) of chloride of silver contain 35.5 (the atomic weight) of chlorine, 8.53 of chloride of silver will be found to contain 2.11 of chlorine:—

$$143.5 : 35.5 :: 8.53 : x$$

$$\begin{array}{r} 8.53 \\ \hline 1.065 \\ 17.75 \\ 284.0 \\ \hline 143.5)302.815(2.11 \\ 287.0 \\ \hline 15.81 \\ 14.35 \\ \hline 1.465 \\ 1.435 \end{array}$$

$$x = 2.11$$

And if 143.5 of chloride of silver contain 108 of silver, 8.53 of chloride of silver will contain 6.42 of silver. To ascertain, for example, the amount of silver in a substance containing, say, nitrate of silver, all that is necessary is to take a weighed quantity of the substance, dissolve it, precipitate the whole of the silver by adding hydrochloric acid or other chloride till more chloride of silver falls, collect the precipitate on a filter, wash, dry, and weigh. The amount of silver in the dried chloride, ascertained by calculation, is the amount of silver in the quantity of substance on which the operation was conducted; a rule-of-three sum gives the quantity per cent.—the form in which the results of quantitative analysis are usually stated. Occasionally a constituent of a substance admits of being isolated and weighed in the uncombined state. Thus the amount of mercury in a substance may be determined by separating and weighing the

mercury in the metallic condition; if occurring as calomel (HgCl) or corrosive sublimate ($HgCl_2$) the proportion of chlorine may then be ascertained by calculation (Hg = 200; Cl = 35.5).

*Nature of Gravimetric Quantitative Analysis.*—As above stated, a body may be isolated and *weighed* and its quantity thus ascertained; or it may be separated and weighed in combination with another body whose combining proportion is well known; this is quantitative analysis by the *gravimetric* method.

*Nature of Volumetric Quantitative Analysis.*—Volumetric operations depend for success on some accurate initial gravimetric operation. A *weighed* amount of a pure salt is dissolved in a given *volume* of water or other fluid and thus forms *a standard solution.* Accurately *measured* quantities of such a solution will obviously contain just as definite amounts of the dissolved salt as if those amounts were actually *weighed* in a balance, and, as measuring occupies less time than weighing, the volumetric operations can be conducted with great economy of time as compared with the corresponding gravimetric operations. Quantitative analysis by the *volumetric* method consists in noting the volume of the standard liquid required to be added to the substance under examination before a given effect is produced. Thus, for instance, a solution of nitrate of silver of known strength may be used in experimentally ascertaining an unknown amount of chlorine in any substance. The silver solution is added to a solution of a definite quantity of the substance until flocks of chloride of silver cease to be precipitated: every 108 parts of silver added (or 170 of nitrate of silver: Ag = 108, N = 14, $O_3$ = 48; total 170) indicates the presence of 35.5 of chlorine, or an equivalent quantity of any chloride. The preparation of standard solutions, such as that of nitrate of silver, to which allusion is here made, requires considerable care; but when made, certain analyses can, as already indicated, be executed with far more rapidity and ease than by gravimetric processes.

*Note.*—The quantitative analysis of solids and liquids often involves determination of temperature and specific gravity. These processes will now be explained, after which an outline of volumetric and gravimetric quantitative analysis will be given. The scope of this work precludes any attempt to describe all the little mechanical details observed by quantitative analysts; essential operations, however, are so fully treated that expert manipulators will meet with little difficulty.

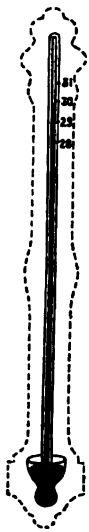
## Measurement of Atmospheric Pressure.

*The Barometer.*—The analysis of gases and vapors involves, also, determinations of the varying pressure of the atmosphere as indicated by the *barometer* (from βάρος, *baros*, weight, and μέτρον, *metron*, measure).

*The ordinary mercurial barometer* is a glass tube 33 or 34 inches long, closed at one end, filled with mercury, and inverted in a small cistern or cup of mercury (fig. 62). The mercury remains in the tube owing to the weight or pressure of the atmosphere on the

exposed surface of the liquid, the average height of the column being nearly 30 inches. In the popular form of the instrument, the wheel-barometer, the cistern is formed by a recurvature of the tube (fig. 63); on the exposed surface of the mercury a float is placed, from which a thread passes over a pulley and moves an index whenever the column of mercury rises or falls. As supplied to the public these barometers are usually inclosed in ornamental frames with thermometers attached. In the wheel-barometer the glass tube and contained column of mercury are altogether inclosed, the index alone being visible. In the other variety the upper end of the glass tube and mercurial column are exposed and the height of the mercury is ascertained by direct observation.

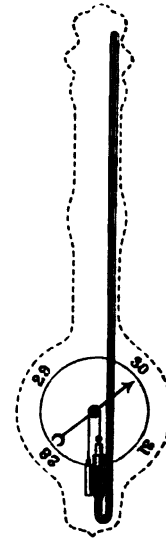
Fig. 62.



Barometer.

*The aneroid barometer* (from ἀ, *a*, *without*, and νηρὸς, *neros*, fluid) consists of a small, shallow, vacuous metal-drum, the sides of which approach each other when an increase of atmospheric pressure occurs, their elasticity enabling them to recede toward their former position on a decrease of pressure. This motion is so multiplied and altered in direction by levers, etc., as to act on a hand traversing a plate on which are marked numbers corresponding with those showing the height of the mercurial column of the ordinary barometer by which the aneroid was adjusted. *The Bourdon barometer* (from the name of the inventor) is a modified aneroid, containing, in the place of the round metal box, a flattened vacuous tube of metal, bent nearly to a circle. These barometers are also useful for measuring the pressure in steam-boilers, etc. Under the name of *pressure-gauges* they are sold to indicate pressure of 500 pounds and upwards per square inch. From their portability (they can be made of 1 to 2 inches in diameter and 1 inch thick) they are excellent companions for travellers wishing to know the height of hills, mountains, and other elevations.

Fig. 63.




Barometer.

For further information concerning the influence of pressure on the volume of a gas or vapor see page 536; and for descriptions of the methods of analyzing gases, refer to Ganot's "Physics" (translated by Atkinson), Miller's "Chemical Physics," and "Analysis of Gases" in Watt's "Dictionary of Chemistry."

## MEASUREMENT OF TEMPERATURE.

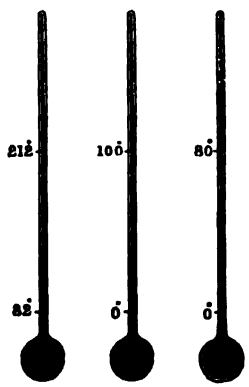
*General Principles.*—As a rule, all bodies expand on the addition, and contract on the abstraction of heat, the alteration in volume being constant and regular for equal increments or decrements of temperature. The extent of this alteration in a given substance, expressed in parts or degrees, constitutes the usual method of intelligibly stating, with accuracy, precision, and minuteness, a particular condition of warmth or temperature, that is, of sensible heat. The substance commonly employed for this purpose is mercury, the chief advantages of which are that it will bear a high temperature without boiling, a low temperature without freezing, does not adhere to glass to a sufficient extent to "wet" the sides of any tube in which it may be inclosed, and, from its good conducting-power for heat, responds rapidly to changes of temperature. Platinum, earthenware, alcohol, and air are also occasionally used for thermometric purposes.

*The Thermometer.*—The construction of an accurate thermometer is a matter of great difficulty; but the following are the leading steps in the operation. Select a piece of glass tubing having a fine capillary (*capillus*, a hair) bore, and about a foot long; heat one extremity in the blowpipe-flame until the orifice closes, and the glass is sufficiently soft to admit of a bulb being blown; heat the bulb to expel air, immediately plunging the open extremity of the tube into mercury; the bulb having cooled, and some mercury having entered and taken the place of expelled air, again heat the bulb and tube until the mercury boils and its vapor escapes through the bore of the tube: again plunge the extremity under mercury, which will probably now completely fill the bulb and tube. When cold the bulb is placed in melting ice. The top of the column of mercury in the capillary tube should then be within an inch or two of the bulb; if higher, some of the mercury must be expelled by heat; if lower, more metal must be introduced as before. The tube is now heated near the open end and a portion drawn out, until the diameter is reduced to about one-tenth. The bulb is next warmed until the mercurial column rises above the constricted part of the tube, which is then rapidly fused in the blowpipe-flame, and the extremity of the tube removed.

The instrument is now ready for *graduation*. The bulb is placed in boiling-water (a medium having, *cæteris paribus*, an invariable temperature), and, when the position of the top of the mercurial column is constant, a mark is made on the tube by a scratching diamond or a file. This operation is repeated with melting ice (also a medium having an invariable tempe-

rature). The space between these two marks is divided into a certain number of intervals termed *degrees*. Unfortunately this number is not uniform in all countries: in England it is 180, as proposed by Fahrenheit; in France 100, as proposed by Celsius (the Centigrade scale), a number generally adopted by scientific men; in some parts of the Continent the divisions are 80 for the same interval, as suggested by Réaumur. Whichever be the number selected, similar markings should be continued beyond the boiling- and freezing-points as far as the length of the stem admits. They may be made on the stem itself, or on any wood, metal, or earthenware frame on which the stem is mounted.

Fig. 64.
Thermometric Scales.




*Thermometric Scales* (fig. 64).—On the Centigrade (C.) and Réaumur (R.) scales the freezing-point of water is made zero, and the boiling-point 100 and 80 respectively; on the Fahrenheit (F.) scale the zero is placed 32 degrees below the congealing-point of water, the boiling-point of which becomes, consequently, 212. Even on the Fahrenheit system temperatures below the freezing-point of water are often spoken of as "degrees of frost;" thus 19 degrees as marked on the thermometer would be regarded as "13 degrees of frost." It is to be regretted that the freezing-point of water is not universally regarded as the zero-point, and the number of intervals between that and the boiling-point everywhere the same.

*The degrees of one scale are easily converted into those of another*, if their relations be remembered, namely: 180 (F.), 100 (C.), 80 (R.); or 18, 10, and 8; or, best, 9, 5, and 4.

*Formulæ for the conversion of degrees of one thermometric scale into those of another.*

F = Fahrenheit. C = Centigrade.
R = Réaumur. D = The observed degree.

If above the freezing-point of water (32° F; 0° C; 0° R),

| | |
|---|---|
| F into C . . . . . | $(D-32)\div 9\times 5$. |
| F " R . . . . . | $(D-32)\div 9\times 4$. |
| C " F . . . . . | $D\div 5\times 9+32$. |
| R " F . . . . . | $D\div 4\times 9+32$. |

If below freezing, but above 0° F (—17°.77 C; —14°.22 R),

F into C . . . . . .—(32—D)÷9×5.
F " R . . . . . .—(32—D)÷9×4.
C " F . . . . . 32—(D÷5×9).
R " F . . . . . 32—(D÷4×9).

If below 0° F (—17°.77 C; —14°.22 R),

F into C . . . . . .—(D+32)÷9 ×5.
F " R . . . . . .—(D+32)÷9 ×4.
C " F . . . . . .—(D÷ 5 ×9)—32.
R " F . . . . . .—(D÷ 4 ×9)—22.

For all degrees:—

C into R . . . . . D÷5×4.
R " C . . . . . D÷4×5.

*In ascertaining the temperature of a liquid*, the bulb of a thermometer is simply inserted and the degree noted. In determining the boiling-point, also, the bulb is inserted in the liquid, if a pure substance. In taking the boiling-point of a liquid which is being distilled from a mixture, the bulb of the thermometer should be near to but not beneath the surface.

*The "boiling-point" of a liquid* is the temperature at which the elasticity of the vapor of the substance overcomes the atmospheric or other pressure to which the liquid is exposed. If the pressure is equal to 760 mm. (29.92 inches) of mercury, water will boil at 100° C. (212° F.). The boiling-point of a drop of a fluid is taken by introducing it into the closed extremity of a small U-tube, the remaining portion of the closed limb being filled with mercury. The tube is lowered into a bath, the open limb being above the surface of the fluid of the bath. The bath is slowly and equally heated, and the boiling-point of the liquid, indicated by the mercury falling until it is level in the two limbs, taken by a thermometer whose bulb is close to the U-tube.

*The following are the boiling-points* of a few substances met with in pharmacy:—

| | Centigrade. | Fahrenheit. |
|---|---|---|
| Alcohol, absolute | 78.3 | 173 |
| " 84 per cent. | 79.5 | 175 |
| " 49 per cent. (proof spirit) | 81.4 | 178.5 |
| " amylic | 132.2 | 270 |
| Benzol | 80.6 | 177 |
| Bromine | 63.0 | 145.4 |
| Benzoic acid | 239.0 | 462 |
| Carbolic acid | 187.8 | 370 |
| Chloroform | 61 | 142 |
| Ether (B. P.) (below) | 40.5 | 105 |
| " pure | 35 | 95 |

| | Centigrade. | Fahrenheit |
|---|---|---|
| Mercury *in vacuo* (as in a thermometer) . | 304 | 580 |
| " in air (barom. at 30 inches) . . | 350 | 662 |
| Water (barom. at 29.92 inches) . . . | 100 | 212 |
| " ( " 29.33 " ) . . . | 99.5 | 211 |
| " ( " 28.74 " ) . . . | 99 | 210 |
| Saturated solutions of— | | |
| Cream of tartar . . . . . | 101 | 214 |
| Common salt . . . . . | 106.6 | 224 |
| Sal ammoniac . . . . . | 113.3 | 236 |
| Nitrate of sodium . . . . . | 119 | 246 |
| Acetate of sodium . . . . . | 124.4 | 256 |
| Chloride of calcium . . . . . | 179.4 | 355 |

By "gentle heat," U. S. P., is meant any temperature between 90° and 100° F.

*To determine Melting-points of Fat.*—Heat a fragment of the substance (spermaceti or wax for example) till it liquefies, and then draw up a small portion into a thin glass tube, about the size of a knitting-needle. Immerse the tube in cold water contained in a beaker, and slowly heat the vessel till the thin opaque cylinder of solid fat melts and becomes transparent; a delicate thermometer placed in the water indicates the point of change to the fifth of a degree. Remove the source of heat, and note the congealing-point of the substance; it will be identical with or close to the melting-point.

*Pyrometers.*—Temperatures above the boiling-point of mercury are determined by ascertaining to what extent a bar of platinum or porcelain has elongated. The bar is inclosed in a cavity of a suitable case, a plug of platinum or porcelain placed at one end of the bar, and the whole exposed in the region the temperature of which is to be found. After cooling, the distance to which the bar has forced the plug along the cavity is accurately measured and the corresponding degree of temperature noted. The value of the distance is fixed for low temperatures by comparison with a mercurial thermometer, and the scale carried upwards through intervals of equivalent length. Such thermometers are conventionally distinguished from ordinary instruments by the name *pyrometer* (from πῦρ, *pūr*, fire, and μέτρον, *metron*, measure).

*The following are melting-points* of substances official in the British Pharmacopœia:—

| | In degrees Centigrade. | In degrees Fahrenheit. |
|---|---|---|
| Acetic acid, glacial | 8.9 | 48 |
| " " " congeals at | 1.1 | 34 |
| Benzoic acid | 120 | 248 |
| Carbolic acid | 35 | 95 |
| Oil of theobroma (about) | 32 | 90 |
| Phosphorus | 43.3 | 110 |
| Prepared lard (about) | 38 | 100 |
| " suet | 39.5 | 103 |
| Spermaceti (not under) | 38 | 100 |
| White wax | 65.5 | 150 |
| Yellow wax | 60 | 140 |

The order of fusibility of a few of the metals is as follows:—

| | In degrees Centigrade. | In degrees Fahrenheit. |
|---|---|---|
| Mercury | — 39.4 | — 39 |
| Potassium | + 62.5 | +144.5 |
| Sodium | 97.6 | 207.7 |
| Tin | 227.8 | 442 |
| Bismuth | 264 | 507 |
| Lead | 325 | 617 |

| | In degrees Centigrade. | In degrees Fahrenheit. |
|---|---|---|
| Zinc | + 411.6 | + 773 |
| Antimony | 621 | 1150 |
| Silver | 1023 | 1873 |
| Copper | 1091 | 1996 |
| Gold | 1102 | 2016 |
| Cast iron | 1530 | 2786 |

## QUESTIONS AND EXERCISES.

922. On what fundamental laws are the operations of quantitative analysis based?

923. What is the general nature of *gravimetric* quantitative analysis?

924. Describe the general principle of *volumetric* quatitative analysis.

925. How are variations in atmospheric pressure quantitatively determined?

926. Explain the construction and mode of action of a mercurial barometer.

927. In what respect does a wheel-barometer differ from an instrument in which the readings are taken from the top of the column of mercury?

928. Describe the principle of action of an aneroid barometer.

929. On what general principles are thermometers constructed?

930. What material is employed in making thermometers?
931. Why is mercury selected as a thermometric indicator?
932. Describe the manufacture of a mercurial thermometer.
933. How are thermometers graduated?
934. Give formulæ for the conversion of the degrees of one thermometric scale into those of another, (*a*) when the temperature is above the freezing-point of water, (*b*) below 32° F. but above 0° F., and (*c*) below 0°.
935 Name the degree C. equivalent to 60° F.
936. What degree C. is represented by —4° F.?
937. Mention the degree F. indicated by 28° C.
938. Convert 100° R. into degrees C. and F.
939. State the boiling-points of alcohol, chloroform, ether, mercury, and water on either thermometric scale.
940. Describe the details of manipulation in estimating the melting-point of fats.
941. In what respect do pyrometers differ from thermometers?
942. Mention the melting-points of glacial acetic acid, oil of theobroma, lard, suet, and wax.
943. Give the fusing-points of tin, lead, zinc, copper, and cast-iron.

## ESTIMATION OF WEIGHT.

### DEFINITIONS.

All bodies, celestial and terrestrial, attract each other, the amount of attraction being in direct proportion to the quantity of matter of which they consist, and in inverse proportion to the squares of their distances. This is *gravitation*. When gravitation in certain directions is exactly counterbalanced by gravitation in opposite directions, a body (*e. g.* the earth) remains suspended in space. Such a body in relation to other bodies, has gravity but not weight. *Weight* is the effect of gravity, being the excess of gravitation in one direction over and above that exerted in the opposite direction. Weight, truly, in any terrestrial substance, is the excess of attraction which it and the earth have for each other over and above the attraction of each in opposite directions by the various heavenly bodies. But practically, the weight of any terrestrial substance is the effect of the attraction of the earth only. *Specific* weight is the definite or precise weight of a body in relation to its bulk; it is more usually but not quite correctly termed *specific gravity*—gravity belonging to the earth, not to the substance.

### QUESTIONS.

944. What is understood by gravitation?
945. State the difference between weight and gravity.
946. Mention a case in which a body has gravity but no apparent weight.
947. Practically, what causes the weight of terrestrial substances?

## Weights and Measures.

*The Balance.*—The balance used in the quantitative operations of analytical chemistry must be accurate and sensitive. The points of suspension of the beam and pans should be polished steel or agate knife-edges, working on agate planes. It should turn easily and quickly, without too much oscillation, to $\frac{1}{10}$ or $\frac{1}{60}$ of a grain, or $\frac{1}{10}$ of a milligramme, when 1000 grains, or 50 or 60 grammes, are placed in each scale. (Grammes are weights of the metric system, a description of which is given on the next two or three pages.) The beam should be light and strong, capable of supporting a load of 1500 grains or 100 grammes; its oscillations are observed by help of a long index attached to its centre, and continued downward for some distance in front of the supporting pillar of the balance. The instrument should be provided with screws for purposes of adjustment, a mechanical contrivance for supporting the beam above its bearing when not in use or during the removal or addition of weights, spirit levels to enable the operator to give it a horizontal position, and be inclosed in a glass case to protect from dust. It should be placed in a room the atmosphere of which is not liable to be contaminated by acid fumes, in a situation free from vibration; and a vessel containing lumps of quicklime should be placed in the case to keep the inclosed air dry and prevent the formation of rust on the steel knife-edges or other parts. During weighing, the doors of the balance should be shut, in order that currents of air may not unequally influence the pans.

*The Weights.*—These should be preserved in a box having a separate compartment for each. They must not be lifted directly with the fingers, but by a small pair of forceps. If grain-weights, they should range from 1000 grs. to $\frac{1}{10}$ gr., a $\frac{1}{10}$ weight being fashioned of gold wire to act as a "rider" on the divided beam, and thus indicate by its position 100ths and 1000ths of a grain. From $\frac{1}{10}$ to 10 grs. the weights may be of platinum; thence upwards to 1000 grs. of brass. The relation of the weights to each other should be decimal. Metric decimal weights may range from 1000 grammes to 1 gramme, of brass, and thence downward to 1 centigramme of platinum, a gold centigramme rider being employed to indicate milligrammes and tenths of a milligramme.

**Weights and Measures of the U. S. Pharmacopœia.**—The weights and measures used by physicians and apothecaries in the United States, when prescribing and preparing medicines, are the following—as stated in the U. S. Pharmacopœia.

*Weights.*—These are derived from the *troy pound*, and are exhibited in the following table, with their signs annexed:—

| | | | |
|---|---|---|---|
| One pound, | ℔ | = 12 ounces | = 5760 grains. |
| One ounce, | ℥ | = 8 drachms | = 480 grains. |
| One drachm, | ʒ | = 3 scruples | = 60 grains. |
| One scruple, | ℈ | . . . . . | = 20 grains. |
| One grain, | gr. | . . . . . | = 1 grain. |

In order to avoid the danger of mistakes from confounding the troy and avoirdupois pounds, the term *pound* is disused in the for-

mulæ of the U. S. Pharmacopœia and the desired weight is expressed in ounces, each containing four hundred and eighty grains. This ounce is always printed *troyounce*, to guard against the error of substituting for it the avoirdupois ounce, consisting of four hundred and thirty-seven and a half grains. The drachm and scruple are also disused, and replaced by their equivalents in grains.

It is highly important that persons engaged in preparing medicines should be provided with troy weights. But those who are not so provided can make their avoirdupois weights available as substitutes for troy weights, by bearing in mind that 42.5 grains, added to the avoirdupois ounce, will make it equal to the troy ounce; and that 1240 grains, deducted from the avoirdupois pound, will reduce it to the troy pound.

*Measures.*—These are derived from the *wine gallon*, and are given in the following table, with their signs annexed:—

| | | | | |
|---|---|---|---|---|
| One gallon, | C | = | 8 pints | = 61,440 minims. |
| One pint, | O | = | 16 fluidounces | = 7,680 minims. |
| One fluidounce, | f℥ | = | 8 fluidrachms | = 480 minims. |
| One fluidrachm, | fℨ | . | . . . . . | = 60 minims. |
| One minim, | ♏ | . | . . . . . | = 1 minim. |

In this work the term *gallon* is not used, that measure being always expressed in pints.

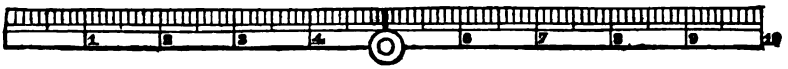
At the temperature of 60°, a pint of distilled water weighs 7291.2 grains, a fluidounce 455.7 grains.

**The Metric System** of weights (the word *metric* is from the Greek *μέτρον*, *metron*, measure) is greatly to be preferred to all others, the relation of the metric weights of all denominations to measures of length, capacity, and surface being so simple as to be within the perfect comprehension of a child; while under the British and American plans the weights have no such relation, either with each other or with the various measures. Moreover the metric system is in perfect harmony with the universal method of counting; it is a decimal system.

[It is perhaps impossible to realize, much more express, the advantages we enjoy from the fact that in every country of the world the system of numeration is identical. That system is the decimal. Whatever language a man speaks, his method of numbering is decimal; his talk concerning number is decimal; his written or printed signs signifying number are decimal. With the figures, 1, 2, 3, 4, 5, 6, 7, 8, 9, 0 he represents all possible variation in number, the position of a figure in reference to its companions alone determining its value, a figure on the left hand of any other figure in an allocation of numeral symbols (for example, 1871) having ten times the value of that figure, while the figure on the right hand of any other has a tenth of the value of that other. When the youngest pupil is asked how many units there are in 1871, he smiles at the simplicity of the question, and says 1871. How many tens? 187, and 1 over. How many hundreds? 18, and 71 over. How many thousands? 1, and 871 over. But if he is asked how many scruples there are in 1871 grains, how many drachms, how many ounces—he first inquires which drachms or which ounces are meant, avoirdupois ounces, troy ounces,

or wine ounces, and then brings out his slate and pencil. And so with the pints or gallons in 1871 fluidounces, or the feet and yards in 1871 inches, or the pence, shillings, and pounds in 1871 farthings; to say nothing of cross questions, such as the value of 1871 articles at 2 dollars and 20 cents per dozen, or of the perplexity caused by the varying values of several individual weights or of measures of length, capacity, and surface in different parts of the country. What is desired is, that there should be an equally simple decimal relation among weights and measures and coins as already universally exists among numbers. This condition of things having already been introduced into most other countries, there is no good reason why it should not be accomplished in the United States and Great Britain.]

Fig. 65.




The Decimetre.

The Metric System of weights and measures is founded on the metre. The engraving (Fig. 65) represents a pocket folding measure, the tenth part of a metre in length, divided into ten centimetres, and each centimetre into 10 millimetres.

The units of the system with their multiples and submultiples are as follows:—

## Units.

*Length.*—The *Unit of Length* is the Metre, derived from the measurement of the Quadrant of a Meridian of the Earth. (Practically it is the length of certain carefully preserved bars of metal, from which copies have been taken.)

*Surface.*—The *Unity of surface* is the Are, which is the Square of Ten Metres.

*Capacity.*—The *Unity of Capacity* is the Litre, which is the Cube of a Tenth Part of a Metre.

*Weight.*—The *Unit of Weight* is the Gram, which is the Weight of that quantity of distilled water, at its maximum density (4° C.) which fills a Cube of the One-hundredth part of the Metre.

TABLE.

*Note.*—Multiples are denoted by the Greek words "Deca," Ten, "Hecto," Hundred, "Kilo," Thousand.
Subdivisions, by the Latin words, "Deci," One-tenth, "Centi," One-hundredth, "Milli," One-thousandth.

| Quantities. | Length. | Surface. | Capacity. | Weight. |
|---|---|---|---|---|
| 1000 | Kilo-metre | . . . | Kilo-litre | Kilo-gram |
| 100 | Hecto-metre | Hectare | Hecto-litre | Hecto-gram |
| 10 | Deca-metre | . . . | Deca-litre | Deca-gram |
| 1(Units) | METRE | ARE | LITRE | GRAM |
| .1 | Deci-metre | . . . | Deci-litre | Deci-gram |
| .01 | Centi-metre | Centiare | Centi-litre | Centi-gram |
| .001 | Milli-metre | . . . | Milli-litre | Milli-gram |

When the metric method is exclusively adopted these Units and Table, comprising the entire System of Weights and Measures, represent all that will be essential to be learned in lieu of the numerous and complicated tables hitherto in use. Adopting the style of elementary books on Arithmetic, the Tables may be expanded in the following manner:—

10 Milligrams make 1 centigram.
10 Centigrams " 1 decigram.
10 Decigrams " 1 gram.
10 Grams " 1 decagram.
10 Decagrams " 1 hectogram.
10 Hectograms " 1 kilogram.

10 Millilitres make 1 centilitre,
etc.

10 Millimetres make 1 centimetre,
etc.

The following approximate equivalents of metrical units should be committed to memory:—

1 Metre = 3 feet 3 inches and 3 eighths.
1 Are = a square whose side is 11 yards.
1 Litre = $1\frac{3}{4}$ pint.
1 Gramme = $15\frac{1}{2}$ grains.

The kilometre is equal to 1100 yards.
The Hectare = $2\frac{1}{2}$ acres nearly.
The Metric Ton of 1000 Kilogrammes = 19 cwt. 2 qrs. 20 lbs. 10 ozs.
The Kilogramme = 2 lbs. $3\frac{1}{4}$ ozs. nearly.

For *exact* equivalents, in many forms, see pages 527 and 528. (The word *gramme* is, in English, usually written *gram*.)

*Decimal Coinage.*—In most countries where the metric system of weights and measures is employed, a decimal division of coins is also adopted. This course, conjoined with the ordinary decimal method of enumerating, which fortunately is in universal use, renders calculations of all kinds most simple—easy to an extent which cannot be conceived in countries like England, where the operations of weigh-

ing, measuring, paying, and counting have only the most absurdly intricate relations to each other.

The General Council under whose authority the British Pharmacopœia is issued encourages medical practitioners and pharmacists in the adoption of the metric system, and gives the annexed statement of metric weights and measures.

## WEIGHTS AND MEASURES OF THE METRICAL SYSTEM.

(From the British Pharmacopœia, of 1867.)

### WEIGHTS.

| | | | |
|---|---|---|---|
| 1 Milligramme | = the thousandth part of one grm. | or 0.001 | grm. |
| 1 Centigramme | = the hundredth " | 0.01 | " |
| 1 Decigramme | = the tenth " | 0.1 | " |
| 1 Gramme | = weight of a cubic centimetre of water at 4° C. | 1.0 | " |
| 1 Decagramme | = ten grammes | 10.0 | " |
| 1 Hectogramme | = one hundred grammes | 100.0 | " |
| 1 Kilogramme | = one thousand grammes | 1000.0 | (1 kilo.). |

### MEASURES OF CAPACITY.

| | | | | | | |
|---|---|---|---|---|---|---|
| 1 Millilitre | = 1 | cub. centim. | or the mea. of | 1 | gram. | of water. |
| 1 Centilitre | = 10 | " | " | 10 | " | " |
| 1 Decilitre | = 100 | " | " | 100 | " | " |
| 1 Litre | = 1000 | " | " | 1000 | " | (1 kilo.). |

### MEASURES OF LENGTH.

| | | | |
|---|---|---|---|
| 1 Millimetre | = the thousandth part of one metre | or 0.001 | metre. |
| 1 Centimetre | = the hundredth " | 0.01 | " |
| 1 Decimetre | = the tenth " | 0.1 | " |
| 1 Metre | = the ten-millionth part of a quarter of the meridian of the earth. | | |

The National Convention for revising the Pharmacopœia of the United States also recognizes the claims of the metric system of weights and measures by giving, in the recent (fifth) edition of the Pharmacopœia, Tables of the units of the metrical system with their multiples and submultiples, similar to the foregoing, and the following Tables showing the relation to each other of the metrical and official or Troy systems.

## RELATION OF WEIGHTS OF THE U. S. PHARMACOPŒIA TO METRICAL WEIGHTS.

| Fraction of a grain in milligrammes. | | | Grains in equivalent metrical weights. | | | Drachms, ounces, and pounds, in equivalent metrical weights. | | |
|---|---|---|---|---|---|---|---|---|
| Grain. | | Milligramme. | Grains. | | Centigrammes. | Drachms. | | Grammes. |
| 1/64 | = | 1.012 | 1 | = | 6.479 | 1 | = | 3.887 |
| 1/60 | = | 1.079 | | | | 2 | = | 7.775 |
| 1/50 | = | 1.295 | | | Decigrammes. | | | |
| 1/48 | = | 1.349 | 2 | = | 1.295 | | | Decagrammes. |
| 1/40 | = | 1.619 | 3 | = | 1.943 | 3 | = | 1.166 |
| 1/36 | = | 1.799 | 4 | = | 2.591 | 4 | = | 1.555 |
| 1/30 | = | 2.159 | 5 | = | 3.239 | 5 | = | 1.943 |
| 1/25 | = | 2.591 | 6 | = | 3.887 | 6 | = | 2.332 |
| 1/24 | = | 2.699 | 7 | = | 4.535 | 7 | = | 2.721 |
| 1/20 | = | 3.239 | 8 | = | 5.183 | | | |
| 1/16 | = | 4.049 | 9 | = | 5.831 | Ounces. | | |
| 1/15 | = | 4.319 | 10 | = | 6.479 | 1 | = | 3.1103 |
| 1/12 | = | 5.399 | 12 | = | 7.775 | 2 | = | 6.2206 |
| 1/10 | = | 6.479 | 15 | = | 9.718 | 3 | = | 9.3309 |
| 1/8 | = | 8.098 | | | | | | |
| 1/6 | = | 10.798 | | | Grammes. | | | Hectogrammes. |
| 1/5 | = | 12.958 | 16 | = | 1.036 | 4 | = | 1.2441 |
| 1/4 | = | 16.197 | 20 | = | 1.295 | 5 | = | 1.5551 |
| 1/3 | = | 21.597 | 24 | = | 1.555 | 6 | = | 1.8661 |
| 1/2 | = | 32 395 | 25 | = | 1.619 | 7 | = | 2.1772 |
| | | | 30 | = | 1.943 | 8 | = | 2.4882 |
| | | | 40 | = | 2.591 | 9 | = | 2.7992 |
| | | | 50 | = | 3.239 | 10 | = | 3.1103 |
| | | | 60 | = | 3.887 | 11 | = | 3.4213 |
| | | | | | | Pounds. | | |
| | | | | | | 1 | = | 3.7324 |
| | | | | | | 2 | = | 7.4648 |
| | | | | | | | | Kilogrammes. |
| | | | | | | 3 | = | 1.1197 |

## RELATION OF METRICAL WEIGHTS TO WEIGHTS OF THE U. S. PHARMACOPŒIA.

| Metrical weights. | | Exact equivalents in grains. | Approximate equivalents in grains. |
|---|---|---|---|
| **Milligrammes.** | | | |
| 1 | = | .0154 | $\frac{1}{65}$ |
| 2 | = | .0308 | $\frac{1}{32}$ |
| 3 | = | .0463 | $\frac{1}{22}$ |
| 4 | = | .0617 | $\frac{1}{16}$ |
| 5 | = | .0771 | $\frac{1}{13}$ |
| 6 | = | .0926 | $\frac{1}{11}$ |
| 7 | = | .1080 | $\frac{1}{9}$ |
| 8 | = | .1234 | $\frac{1}{8}$ |
| 9 | = | .1389 | $\frac{1}{7}$ |
| **Centigrammes.** | | | |
| 1 | = | .1543 | $\frac{1}{6}$ |
| 2 | = | .3086 | $\frac{1}{3}$ |
| 3 | = | .4630 | $\frac{1}{2}$ |
| 4 | = | .6173 | $\frac{2}{3}$ |
| 5 | = | .7717 | $\frac{3}{4}$ |
| 6 | = | .9260 | $\frac{9}{10}$ |
| 7 | = | 1.0803 | 1 |
| 8 | = | 1.2347 | $1\frac{1}{4}$ |
| 9 | = | 1.3890 | $1\frac{1}{3}$ |
| **Decigrammes.** | | | |
| 1 | = | 1.543 | $1\frac{1}{2}$ |
| 2 | = | 3.086 | 3 |
| 3 | = | 4.630 | $4\frac{1}{2}$ |
| 4 | = | 6.173 | 6 |
| 5 | = | 7.717 | $7\frac{1}{2}$ |
| 6 | = | 9.260 | 9 |
| 7 | = | 10.803 | 11 |
| 8 | = | 12.347 | $12\frac{1}{2}$ |
| 9 | = | 13.890 | 14 |

| Metrical weights. | | Exact equivalents in grains. | Approximate equivalents in Troy weight. |
|---|---|---|---|
| **Grammes.** | | | |
| 1 | = | 15.434 | gr. xv. |
| 2 | = | 30.868 | ʒss. |
| 3 | = | 46.302 | ℈ij |
| 4 | = | 61.736 | ʒi. |
| 5 | = | 77.170 | ℈iv. |
| 6 | = | 92.674 | ʒiss. |
| 7 | = | 108.038 | ℈vss. |
| 8 | = | 123.472 | ʒij. |
| 9 | = | 138.906 | ℈vij. |
| **Decagrammes.** | | | |
| 1 | = | 154.340 | ʒiiss. |
| 2 | = | 308.680 | ʒv. |
| 3 | = | 463.020 | ʒviiss. |
| 4 | = | 617.360 | ʒx. |
| 5 | = | 771.701 | ʒxiij. |
| 6 | = | 926.041 | ʒxv. |
| 7 | = | 1,030.382 | ʒxviij. |
| 8 | = | 1,234.721 | ʒxx. |
| 9 | = | 1,389.062 | ʒxxiij. |
| **Hectogrammes.** | | | |
| 1 | = | 1,543.402 | ʒiij ℈v. |
| 2 | = | 3,086.804 | ʒvj ʒiij. |
| 3 | = | 4,630.206 | ℥ix ʒv. |
| 4 | = | 6,173.609 | ℔i ʒvij |
| 5 | = | 7,717.011 | ℔i ʒiv |
| 6 | = | 9,260.413 | ℔i ʒvij |
| 7 | = | 10,803.816 | ℔i ʒx ʒiv. |
| 8 | = | 12,347.218 | ℔ij ℥j ʒv. |
| 9 | = | 13,890.620 | ℔ij ʒv. |
| **Kilogramme.** | | | |
| 1 | = | 15,434.023 | ℔ij ʒviij. |
| **Myriogramme.** | | | |
| 1 | = | 154,340.23 | ℔xxvi. ℥ix ʒiv. |

The following Tables, from the British Pharmacopœia, the United States Pharmacopœia, and the Diary of Messrs. De La Rue, will be found useful for reference:—

## WEIGHTS AND MEASURES OF THE BRITISH PHARMACOPŒIA OF 1867.

### WEIGHTS.

| | | |
|---|---|---|
| 1 Grain | gr. | |
| 1 Ounce | oz. | = 437.5 grains. |
| 1 Pound | lb. = 16 ounces | = 7000 " |

### MEASURES OF CAPACITY.

| | | | |
|---|---|---|---|
| 1 Minim | min. | | |
| 1 Fluidrachm | fl. dr. | = | 60 minims. |
| 1 Fluidounce | fl. oz. | = | 8 fluidrachms. |
| 1 Pint | O. | = | 20 fluidounces. |
| 1 Gallon | C. | = | 8 pints. |

### MEASURES OF LENGTH.

1 line = $\frac{1}{12}$ inch.

1 inch = $\frac{1}{39.1393}$ seconds-pendulum.

12 " = 1 foot.
36 " = 3 feet = 1 yard.

Length of pendulum vibrating seconds of mean time in the latitude of London in a vacuum at the level of the sea . . . . . } 39.1393 inches.

(1 cubic inch of distilled water at 62° F. and 30 inches Barom. = 252.458 grains.)

### RELATION OF MEASURES TO WEIGHTS.

| | | | | |
|---|---|---|---|---|
| 1 Minim | is the measure of | | 0.91 | grain of water. |
| 1 Fluidrachm | " | | 54.68 | grains of water. |
| 1 Fluidounce | " | 1 ounce or | 437.5 | " |
| 1 Pint | " | 1.25 pound or | 8750.0 | " |
| 1 Gallon | " | 10 pounds or | 70,000.0 | " |

(Gtt. = *guttæ*, drops. The term "drop" indicates a quantity which is indefinite, and should only be used when approximativeness is alone desired.)

### RELATION OF WEIGHTS AND MEASURES OF THE U. S. PHARMACOPŒIA TO EACH OTHER.

In distilled water at a temperature at 60°.

| | | | | | |
|---|---|---|---|---|---|
| One Pound | = | 0.7900031 | Pint | = | 6067.2238 Minims |
| One Ounce | = | 1.0533376 | Fluidounce | = | 505.6019 Minims. |
| One Drachm | = | 1.0533376 | Fluidrachm | = | 63.2002 Minims. |
| One Scruple | | . . . . . | | = | 21.0667 Minims. |
| One Grain | | . . . . . | | = | 1.0533 Minim. |
| One Gallon | = | 10.1265427 | Pounds | = | 58328.8862 Grains. |
| One Pint | = | 1.2658178 | Pound | = | 7291.1107 Grains. |
| One Fluidounce | = | 0.9493633 | Ounce | = | 455.6944 Grains. |
| One Fluidrachm | = | 0.9493633 | Drachm | = | 56.9618 Grains. |
| One Minim | | . . . . . | | = | 0.9493 Grain. |

### RELATION OF MEASURES OF THE U. S. PHARMACOPŒIA TO CUBIC MEASURE.

| | | | |
|---|---|---|---|
| One Gallon | = | 231. | Cubic Inches. |
| One Pint | = | 28.875 | Cubic Inches. |
| One Fluidounce | = | 1.80468 | Cubic Inch. |
| One Fluidrachm | = | 0.22558 | Cubic Inch. |
| One Minim | = | 0.00375 | Cubic Inch. |

## Metrical Measures of Length.

| | In English inches. | In English feet = 12 inches. | In English yards = 3 feet. | In English fathoms = 6 feet. | In English miles = 1760 yards. |
|---|---|---|---|---|---|
| Millimetre . . . . | 0.03937 | 0.003281 | 0.0010936 | 0.0005468 | 0.0000006 |
| Centimetre . . . . | 0.39371 | 0.032809 | 0.0109363 | 0.0054682 | 0.0000062 |
| Decimetre . . . . | 3.93708 | 0.328090 | 0.1093633 | 0.0546816 | 0.0000621 |
| **Metre** . . . . | **39.37079** | 3.280899 | 1.0936331 | 0.5468165 | 0.0006214 |
| Decametre . . . . | 393.70790 | 32.808992 | 10.9363310 | 5.4681655 | 0.0062138 |
| Hectometre . . . . | 3937.07900 | 328.089920 | 109.3633100 | 54.6816550 | 0.0621382 |
| Kilometre . . . . | 39370.79000 | 3280.899200 | 1093.6331000 | 546.8165500 | 0.6213824 |
| Myriometre . . . . | 393707.90000 | 32808.992000 | 10936.3310000 | 5468.1655000 | 6.2138244 |

1 inch = 2.539954 centimetres.
1 foot = 3.0479449 decimetres.
1 yard = 0.9143835 metre.
1 mile = 1.6093149 kilometre.

## Metrical Measures of Surface.

| | In English square feet. | In Eng. sq yards = 9 square feet. | In English poles = 272.25 sq. feet. | In English roods = 10890 sq. feet. | In English acres = 43560 sq. feet. |
|---|---|---|---|---|---|
| Centiare or square metre . . | 10.764299 | 1.196033 | 0.0395383 | 0.0009885 | 0.0002471 |
| Are or 100 square metres . . | 1076.429934 | 119.603326 | 3.9538290 | 0.0988457 | 0.0247114 |
| Hectare or 10,000 square metres | 107642.993418 | 11960.332602 | 395.3828959 | 9.8845724 | 2.4711431 |

1 square inch = 6.4513669 square centimetres.
1 square foot = 9.2899683 square decimetres.
1 square yard = 0.83609715 square metre or centiare.
1 acre = 0.40467102 hectare.

## Metrical Measures of Capacity.

| | In cubic inches. | In cubic feet = 1728 cubic inches. | In pints = 34.65923 cubic inches. | In gallons = 8 pints = 277.27384 cubic inches | In bushels=8 gallons = 2218.19072 cubic inches. |
|---|---|---|---|---|---|
| Millilitre, or cubic centimetre . | 0.06103 | 0.000035 | 0.00176 | 0.0002201 | 0.0000275 |
| Centilitre, or 10 cubic centimetres | 0.61027 | 0.000353 | 0.01761 | 0.0022010 | 0.0002751 |
| Decilitre, or 100 cubic centimetres | 6.10271 | 0.003532 | 0.17608 | 0.0220097 | 0.0027512 |
| **Litre**, or cubic decimetre . . | 61.02705 | 0.035317 | **1.76077** | 0.2200967 | 0.0275121 |
| Decalitre, or centistere . . | 610.27052 | 0.353166 | 17.60773 | 2.2009668 | 0.2751208 |
| Hectolitre, or decistere . . | 6102.70515 | 3.531658 | 176.07734 | 22.0096677 | 2.7512085 |
| Kilolitre, or stere, or cubic metre | 61027.05152 | 35.316581 | 1760.77341 | 220.0966767 | 27.5120846 |
| Myriolitre, or decastere . . | 610270.51519 | 353.165807 | 17607.73414 | 2200.9667675 | 275.1208459 |

1 cubic inch = 16.386176 cubic centimetres. 1 cubic foot = 28.315312 cubic decimetres. 1 gallon = 4.535926 litres.

## Metrical Measures of Weight.

| | In English grains. | In troy ounces = 480 grains. | In avoirdupois lbs. = 7000 grains. | In cwts. = 112 lbs. = 784000 grains. | Tons = 20 cwts. = 15680000 grains. |
|---|---|---|---|---|---|
| Milligramme . . . . . | 0.01543 | 0.000032 | 0.0000022 | 0.0000000 | 0.0000000 |
| Centigramme . . . . . | 0.15432 | 0.000322 | 0.0000220 | 0.0000002 | 0.0000000 |
| Decigramme . . . . . | 1.54323 | 0.003215 | 0.0002205 | 0.0000020 | 0.0000001 |
| **Gramme** . . . . . | **15.43235** | 0.032151 | 0.0022046 | 0.0000197 | 0.0000010 |
| Decagramme . . . . . | 154.32349 | 0.321507 | 0.0220462 | 0.0001968 | 0.0000098 |
| Hectogramme . . . . . | 1543.23488 | 3.215073 | 0.2204621 | 0.0019684 | 0.0000984 |
| Kilogramme . . . . . | 15432.34880 | 32.150727 | 2.2046213 | 0.0196841 | 0.0009842 |
| Myriogramme . . . . . | 154323.48800 | 321.507267 | 22.0462126 | 0.1968412 | 0.0098421 |

1 grain = 0.064799 gram. 1 troy oz. = 31.103496 grams. 1 lb. avd. = 0.453593 kilogr. 1 cwt. = 50.802377 kilogrs.

## QUESTIONS AND EXERCISES.

948. Mention some advantages of a decimal system of weights and measures.
949. What is the name of the chief unit of the metric decimal system of weights and measures?
950. Mention the names of the metric units of surface, capacity, and weight, and state how they are derived from the unit of length.
951. How are multiples of metric units indicated?
952. State the designations of submultiples of metric units.
953, How many metres are there in a kilometre?
954. How many millimetres in a metre?
955. How many grams in 5 kilograms?
856. How many milligrams in 13½ grams?
957. In 1869 centigrams how many grams?
958. In a metre measure 5 centimetres wide and 1 centimetre thick, how many cubic centimetres?
959. How many litres are contained in a cubic metre of any liquid?
960. State the British equivalent of the metre.
961. How many square yards in an are?
962. How many fluidounces in a litre?
963. How many ounces in a kilogram?
964. Give the relation of a metric ton (1000 kilos.) to a British ton.
965. How many grains are there in 1 ton?
966. How many ounces in 1 ton?
967. How many grains of water in 1 fluidrachm?
968. How many minims in 1 pint?
969. How many grains in 1 pint of water?
970. Whence is the British unit of length derived?

---

## Specific Weight or Specific Gravity.

The *specific weight* of a substance is its weight in comparison with weights of similar bulks of other substances. This comparative heaviness of *solids and liquids* is conventionally expressed in relation to water: they are considered as being lighter or heavier than water. Thus, water being regarded as unity = 1, the relative weight, or specific weight, of ether is represented by the figures .720 (it is nearly three-fourths, .750, the weight of water), oil of vitriol by 1.843 (it is nearly twice, 2.000, as heavy as water). The specific weight of substances is, moreover, the weight of similar volumes *at sixty degrees* (60° F.); for the weight of a definite volume of any substance will vary according to temperature, becoming heavier when cooled, and lighter when heated, different bodies (gases excepted) differing in their rate of contraction and expansion. While, then, specific weight, or, conventionally, *specific gravity*, is truly the comparative weight of equal bulks, the numbers which in Ame-

rica and Great Britain commonly represent specific gravities are the comparative weights of equal bulks at 60° F., water being taken as unity.* The standard of comparison for gases was formerly air, but is now usually hydrogen.

## Specific Gravity of Liquids.

Procure any small bottle holding from 100 to 1000 grains (fig. 66) and having a narrow neck; counterpoise it in a delicate balance; fill it to about halfway up the neck with pure distilled water having a temperature of 60° F.: ascertain the weight of the water, and, for convenience, add or subtract a drop or two, so that the weight shall be a round number of grains; mark the neck by a diamond or file-point at the part cut by the lower edge of the curved surface of the water. Consecutively fill up the bottle to the neck-mark with several other liquids, cooled or warmed to 60° F., first rinsing out the bottle once or twice with a small quantity of each liquid, and note the weights; the respective figures will represent the relative weights of equal bulks of the liquids. If the capacity of the bottle is 10, 100, or 1000 grains, the resulting weights will, without calculation, show the specific gravities of the liquids;

Fig. 66. Fig. 67. Fig. 68. Fig. 69.

Specific-Gravity Bottles.

if any other number, a rule-of-three sum must be worked out to ascertain the weight of the liquids as compared with 1 (or

* The true weight of the body is its weight in air plus the weight of an equal bulk of air and minus the weight of a bulk of air equal to the bulk of brass or other weights employed; or, in other words, its weight *in vacuo* uninfluenced by the buoyancy of the air; but such a correction of the weight of a body is seldom necessary or, indeed, desirable. *Density* is sometimes improperly regarded as synoymous with *specific gravity*. It is true that the density of a body is in exact proportion to its specific gravity; but the former is more correctly the comparative bulk of equal weights, while specific gravity is the comparative weight of equal bulks.

1.000) of water. Bottles conveniently adjusted to contain 250, 500, or 1000 grains, or 100 or 50 grammes of water, when filled to the top of their perforated stopper (Fig. 68), and other forms of the instrument (Fig. 67 and 69), are sold by all chemical apparatus makers. Figure 69 is that of a bottle extremely useful in ascertaining the specific gravities of very volatile liquids.

Verify some of the following stated specific gravities of official liquids:—

| | |
|---|---|
| Acid, acetic, B. P. | 1.044 |
| " " U. S. P. | 1.047 |
| " " diluted, B. P. and U. S. P. | 1.006 |
| " " glacial | 1.065 to 1.066 |
| " carbolic | 1.065 |
| " hydriodic, diluted | 1.112 |
| " hydrochloric, B. P. and U. S. P. | 1.160 |
| " " diluted, B. P. | 1.052 |
| " " " U. S. P. | 1.038 |
| " hydrocyanic, B. P. and U. S. P. | 0.997 |
| " lactic, U. S. P. | 1.212 |
| " nitric, B. P. and U. S. P. | 1.420 |
| " " diluted, B. P. | 1.101 |
| " " " U. S. P. | 1.068 |
| " nitro-hydrochloric | 1.074 |
| " phosphoric, diluted, B. P. | 1.080 |
| " " " U. S. P. | 1.056 |
| " sulphuric, B. P. and U. S. P. | 1.843 |
| " " aromatic | 0.927 |
| " " diluted, B. P. | 1.094 |
| " " U. S. P. | 1.082 |
| " sulphurous, solution of, B. P. | 1.040 |
| " " " U. S. P. | 1.035 |
| Alcohol, U. S. P. | 0.835 |
| " absolute | 0.795 |
| " (rectified spirit, 84 per cent.) | 0.838 |
| " (proof spirit, 49 per cent.) | 0.920 |
| " dilutum, U. S. P. | 0.941 |
| " fortius, U. S. P. | 0.817 |
| " amylic, B. P. and U. S. P. | 0.818 |
| Ammonia, aromatic spirit of, B. P. | 0.870 |
| " stronger water of, U. S. P. | 0.900 |
| " solution of, B. P. | 0.959 |
| " strong solution of, B. P. | 0.891 |
| Antimony, solution of chloride of, B. P. | 1.470 |
| Arsenic, hydrochloric solution of, B. P. | 1.009 |
| Arsenical solution (*Liquor Arsenicalis*), B. P. | 1.009 |
| Benzol, B. P. | 0.850 |
| Bismuth and ammonia, solution of citrate of, B. P. | 1.122 |
| Bromine | 2.966 |

| | |
|---|---|
| Chlorine, solution of, B P. | 1.003 |
| Chloroform, B. P. and U. S. P. | 1.490 |
| " spirits of, B. P. | 0.871 |
| Cinchona, liquid extract of yellow, B. P. (about) | 1.100 |
| Creasote, U. S. P. | 1.046 |
| " | 1.071 |
| Ether, B. P. | 0.735 |
| " U. S. P. | 0 750 |
| " pure, B. P. | 0.720 |
| " fortior, U. S. P. | 0.728 |
| Glycerine, B. P. and U. S. P. | 1.250 |
| Iron, solution of pernitrate of, B. P. | 1.107 |
| " " " " " U. S. P. | 1.065 |
| " " " persulphate of, B. P. | 1 441 |
| " " " " " U. S. P. | 1.320 |
| " strong solution of perchloride of, B. P. | 1.440 |
| " tincture of perchloride of, B. P. and U. S. P | 0.992 |
| Lead, solution of subacetate of, B. P. | 1.260 |
| " " " " " U. S. P. | 1.267 |
| Lime, saccharated solution of, B. P. | 1.052 |
| " solution of chlorinated, B. P. | 1.035 |
| Mercury (at 0° C. = 32° F.) | 13.596 |
| " (at 15°.55 C. = 60° F.) | 13.560 |
| " acid solution of nitrate of | 2.246 |
| " " " " " U. S. P. | 2.165 |
| Nitre, sweet spirit of | 0.845 |
| " " " " U. S. P. | 0.837 |
| Oil of mustard, B. P. | 1.015 |
| Potash, solution of, B. P. | 1.058 |
| " " " U. S. P. | 1.065 |
| Soda, " " B. P. | 1.047 |
| " " " U. S P. | 1.071 |
| " " " chlorinated, B. P. | 1.103 |
| " " " " U. S. P. | 1.045 |
| Squill, oxymel of, B. P. | 1.320 |
| Syrup, B. P. | 1.330 |
| " U. S. P. | 1.317 |
| " of buckthorn, B. P. | 1.320 |
| " of chloral | 1.218 |
| " of ginger | |
| " of hemidesmus, B. P. | 1.335 |
| " of iodide of iron, B. P. | 1.385 |
| " of lemons, B. P. | 1.340 |
| " of mulberries, B. P. | 1.330 |
| " of orange-flower, B. P. | 1.330 |
| " of orange-peel, B. P. | |
| " of phosphate of iron, B. P. | |
| " of poppies, B. P. | 1.320 |
| " of red poppy, B. P. | 1.330 |
| " of red roses, B. P. | 1.335 |
| " of rhubarb, B. P. | |

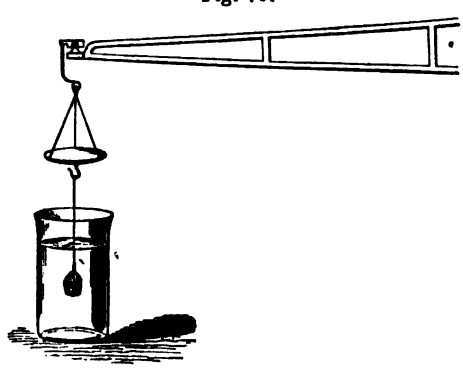
| | | |
|---|---|---|
| Syrup of senna, B. P. | | 1.310 |
| " of squill, B. P. | | |
| " of tolu, B. P. | | 1.330 |
| Treacle, B. P. | (about) | 1.400 |

*Hydrometers.*—The specific gravity of liquids may be ascertained, without scales and weights, by means of an *hydrometer*—an instrument usually of glass, having a graduated stem and a bulb or bulbs at the lower part. The specific gravity of a liquid is indicated by the depth to which the hydrometer sinks in the liquid, the zero of the scale marking the depth to which it sinks in pure water. Hydrometers constructed for special purposes are known under the names of saccharometer, galactometer, elæometer, urinometer, alcoholometer. Hydrometers require a considerable quantity of liquid to fairly float them, and specific gravities observed with them are less delicate and trustworthy than those obtained by the balance, nevertheless they are exceedingly useful for many practical purposes where the employment of a delicate balance would be inadmissible.

## Specific Gravity of Solids in Mass.

Weigh a piece (50 to 250 grains) of any solid substance heavier than water in the usual manner. Then weigh it in water, by suspending it from a shortened balance pan by a fine thread or hair and immersing in a vessel of water (Fig. 70).

Fig. 70.



Weighing a Solid in Water.

The buoyant properties of the water will cause the solid to apparently lose weight; *this loss in weight is the exact weight of an equal bulk of water.* The weight of the substance and the weight of an equal bulk of water being thus ascertained, a rule-of-three sum shows the proportional weight of the substance

to 1.000 of water. To express the same thing by rule, divide the weight in air by the loss of weight in water, the resulting number is the specific gravity in relation to 1 part of water, the conventional standard of comparison.

Verify some of the following specific gravities:—

| | |
|---|---|
| Aluminium | 2.56 |
| Antimony | 6.71 |
| Bismuth | 9.83 |
| Coins, English, gold | 17.69 |
| " " silver | 10.30 |
| " " bronze | 8.70 |
| Copper | 8.95 |
| Gold | 19.34 |
| Iron | 7.84 |
| Lead | 11.36 |
| Magnesium | 1.74 |
| Marble | 2.70 |
| Phosphorus | 1.77 |
| Platinum | 21.53 |
| Silver | 10.53 |
| Sulphur | 2.05 |
| Tin | 7.29 |
| Zinc | 7.14 |

Specific gravities of solid substances should be taken in water having a temperature of about 60° F. The body should be immersed about half an inch below the surface of the water; adhering air-bubbles must be carefully removed; the body must be quite insoluble in water.

### Specific Gravity of Solids in Powder or Small Fragments.

Weigh the particles; place them in a counterpoised specific-gravity bottle of known capacity, and fill up with water, taking care that the substance is thoroughly wetted; again weigh. From the combined weights of water and substance subtract the amount due to the substance; the residue is the weight of the water. Subtract this weight of water from the quantity which the bottle normally contains; the residue is the amount of water displaced by the substance. Having thus obtained the weights of equal bulks of water and substance, a rule-of-three sum shows the relation of the weight of the substance to 1 part of water, the specific gravity.

Or, suspend a cup, short glass tube, or bucket from a shortened balance-pan; immerse in water; counterpoise; place the weighed powder in the cup, and proceed as directed for taking the specific gravity of a solid in mass.

This operation may be conducted on fragments of any of the substances the specific gravities of which are given in the foregoing Table, or on a powdered piece of marble the specific gravity of which has been taken in mass. The specific gravity of one piece of glass, first in mass then in powder, may be ascertained; the result should be identical. The specific gravity of shot is about 11.350; sand, 2.600; mercury, 13.56.

### Specific Gravity of Solids Soluble in Water.

Weigh a piece of sugar or other substance soluble in water; suspend it from a balance in the usual manner, and weigh it in turpentine, benzol, or petroleum, the specific gravity of which is known or has been previously determined; the loss in weight is the weight of an equal bulk of the turpentine. Ascertain the weight of an equal bulk of water by calculation.

| Sp. gr. of turpentine | : | sp. gr. of water | : : | observed bulk of turp. | : | equal bulk of water. |
|---|---|---|---|---|---|---|

The exact weights of equal bulks of sugar and water being obtained, the weight of a bulk of sugar corresponding to 1.000 of water is shown by a rule-of-three sum; in other words, divide the weight of sugar by that of the equal bulk of water, the quotient is the specific gravity of sugar. The stated specific gravity of the sugar ranges from 1.590 to 1.607.

### Specific Gravity of Solids Lighter than Water.

This is obtained in a manner similar to that for solids heavier than water; but the light body is sunk by help of a piece of heavy metal, the bulk of water which the latter displaces being deducted from the bulk displaced by both; the remainder is the weight of a bulk of water equal to the bulk of the light body. For instance, a piece of wood weighing 12 grammes (or grains) is tied to a piece of metal weighing 22 grammes, the loss of weight of the metal in water having been previously found to be 3 grammes. The two, weighing 34 grammes, are now immersed, and the loss in weight found to be 26 grammes. But of this loss 3 grammes have been proved to be due to the buoyant action of the water on the lead; the remaining 23, therefore, represent the same effect on the wood; 23 and 12, therefore, represent the weights of equal bulks of water and wood. As 23 are to 12 so is 1 to .5217. Or, shortly, as before, divide the weight in air by the weight of an equal bulk of water; .5217 is the specific gravity of the wood.

Another specimen of wood may be found to be three-fourths (.750) the weight of water, and others heavier. Cork varies from .100 to .300.

### Specific Gravity of Gases.

This operation is similar to that for liquids. A globe exhausted of air and holding from 1 to 4 litres (or quarts) is suspended from the arm of a balance, and counterpoised by a similar flask. Gases are introduced in succession and their weights noted. A rule-of-three sum shows their specific gravity in relation to air or hydrogen, whichever be taken as a standard.

*Correction of the volume of Gases for Pressure.*—The height of the barometer at the time of manipulation is noted. Remembering the fact that "the bulk of a gas is inversely as the pressure to which it is subjected" (Boyle and Mariotte), a simple calculation shows the volume which the gas would occupy at 760 millimetres (or 29.922 inches), the standard pressure. (30 inches is sometimes adopted as the standard in England.*) Thus, 40 volumes of a gas at 740 millimetres pressure are reduced to 39 when the pressure becomes 760 millimetres (or 90 vols. at 29 ins. barom. become 87 vols. at 30 inches.)

*Correction of the Volume of Gases for Temperature.*—This is done in order to ascertain what volume the gas would occupy at 0° C. (32° F.) or 15°.5 C. (60° F.), according to the standard taken. Gases are equally affected by equal variations in temperature (Charles). They expand about 0.3665† per cent. ($\frac{1}{273}$) of their volume *at the freezing point of water* for every C. degree (0.2036, or $\frac{1}{491}$ for every F. degree) (Regnault). Thus 8 volumes of gas at 0° C. will become 8.293 at 10° C.; for if 100 become 103.665 on being increased in temperature 10° C., 8 will become 8.293 (or if 100 become 102.036 on being increased 10° F., 8 will become 8.1629).

*Vapor-density.*—Vapors are those gases which condense to liquids at common temperatures. By the density of a vapor is meant its specific gravity. The density of a vapor is the ratio of any given volume to a similar volume of air or hydrogen at the same temperature and pressure. But, for convenience of comparison, this experimental specific gravity is referred, by calculation as just described

---

* In France the conventional standard height of the barometer is 760 millimetres at 0° C. (32° F.); in England it is 30 inches, the temperature of the mercurial column being 60° F. 760 millims. is equivalent to 29.922 inches; but the expansion of the metal between 32° F. and 60° F. increases the length of the column to 30.005 inches. The standards are, therefore, almost identical, difference in true length being counterbalanced by the temperature at which the length is observed.

† Corrected for the difference between the mercurial and air thermometers the coefficient of expansion of air is 0.003656 (Miller). The coefficient of expansion of different gases varies very slightly, being somewhat higher for the more liquefiable gases.

for permanent gases, to a temperature of 0° C., and 760 millimetres barom. A teaspoonful or so of liquid is placed in a weighed flask of about the capacity of a common tumbler and having a capillary neck; the flask is heated in an oil-bath to a temperature considerably above the boiling-point of the liquid; at the moment vapor ceases to escape, the neck is sealed by a blowpipe-flame, and the temperature of the bath noted; the flask is then removed, cooled, cleaned, and weighed; the height of the barometer is also taken. The neck of the flask is next broken off beneath the surface of water or mercury (which rush in and fill it), and again weighed, by which its capacity in cub. centims. is found. From these data the volume of vapor yielded by a given weight of liquid is ascertained by a few obvious calculations. The capacity of the globe having been ascertained, the weight of an equal bulk of air* is obtained by a rule-of-three sum. This weight of air is deducted from the original weight of the flask, which gives the true weight of the glass. The weight of the glass is next subtracted from the weight of the flask and contained vapor (now condensed), which gives the weight of material used in the experiment. The volume which this weight of material occupied at the time of experiment is next corrected for temperature (to 0° C.) and pressure (760 millimetres) in the manner just described. The weight of a similar volume of hydrogen is next found.† The weights of equal volumes of hydrogen and vapor being thus determined, the amount of vapor corresponding to one of hydrogen (the specific gravity or vapor-density) is shown by a short calculation. This process of finding the weight of a given volume of vapor is by Dumas. Gay-Lussac's consists in determining the volume of a given weight, it has been improved by Hofmann.

Experiment shows that the specific gravities of many gases and vapors on the hydrogen scale and the proportions in which they combine by weight are identical. Thus, chlorine is 35.5 times as heavy as hydrogen, and 35.5 parts unite with 1 of hydrogen to form hydrochloric acid gas. Hence, if the specific gravity of a gas or vapor is known, its combining proportion may be predicated with reasonable certainty, and *vice versâ*. In applying this rule to gaseous or vaporous compounds, attention must be paid to the extent to which their constituent gases contract at the moment of combination or expand at the moment of decomposition. Thus, steam is found to be com-

---

* 1 cub. centim. of air at 0° C. and 760 millims. weighs 0.001293 gramme.

† 1 litre (1000 cub. centims.) of hydrogen at 0° C. and 760 millimetres (the barometer being at 0° C.) weighs 0.0896 gramme—a volume sometimes termed a *crith* (from κριθή, *krithē*, a barley-corn—figuratively, a small weight); thus a litre of oxygen weighs 16 criths, chlorine 35.5 criths, etc. 100 cubic inches of hydrogen at 32° F. weigh 2.265 grains; at 60° F. 2.143 grains (the barometer being 30 ins. at 60° F. in both cases). 100 cubic inches of air at 32° F. weigh 32.698 grains; at 60° F., 30.935 (barom. 30 ins. at 60° F.). 1 cubic inch of water weighs 252.5 (252.458 at 62° F., and 30 ins. bar.) grains. 1 gallon of water contains 277¼ (277.274 at 62° F.) cubic inches.

posed of two volumes of hydrogen and one of oxygen, the three volumes of constituents condensing to two at the moment of combination. Hence, steam may be expected to be nine times as heavy as hydrogen, which experiment confirms.

These relations may be so expressed as to include both elementary and compound gases and vapors, thus: *molecular weights and specific weights are identical.* Molecular weights represent two volumes of a gas; specific gravity conventionally represents the relative weight of a gas compared with *one* volume of hydrogen or air; hence the specific gravity of a gas or vapor on the H scale is found by calculation on simply dividing the molecular weight by 2; on the air-scale by dividing the hydrogen numbers by 14.44. For example,—

| Name. | Molecular formula. | Molecular weight. | Specific gravity. H=2. | H=1. | Air=1. |
|---|---|---|---|---|---|
| Hydrogen, | $H_2$ | 2 | 2 | 1 | .069 |
| Chlorine, | $Cl_2$ | 71 | 71 | 35.5 | 2.460 |
| Oxygen, | $O_2$ | 32 | 32 | 16 | 1.108 |
| Nitrogen, | $N_2$ | 28 | 28 | 14 | .970 |
| Steam, | $H_2O$ | 18 | 18 | 9 | .625 |
| Ammonia-gas, | $NH_3$ | 17 | 17 | 8.5 | .589 |
| Carbonic acid gas, | $CO_2$ | 44 | 44 | 22 | 1.524 |
| Alcohol (vapor), | $C_2H_6O$ | 46 | 46 | 23 | 1.593 |
| Air, | ..... | ..... | 28.88 | 14.44 | 1.000 |

These specific gravities closely correspond with those obtained by actual experiment. The specific gravity of any gas or vapor may therefore be calculated if the following data are at hand: (*a*) formula, (*b*) atomic weight of constituent elements; these give the molecular weight, and *the molecular weight divided by* 2 *is the specific gravity* on the hydrogen-scale. Specific gravity on the air-scale is then deducible, if (*c*) the specific gravity of air (14.44) in relation to hydrogen be remembered. The absolute weight of any volume of a gas or vapor on the metric system is then obtainable if (*d*) the weight of a litre of hydrogen (0.0896 gramme) be known, or on the English plan by remembering (*e*) that 100 cubic inches of hydrogen at 60° F. weigh 2.143 grains (100 cubic inches of air at 60° F. weigh 30.935 grains).

In confirmation of these statements regarding the mutual relation of specific gravity and atomic weight, a remarkable fact may be mentioned. Regnault several years ago found the weights of 1 litre of hydrogen and oxygen to be respectively .089578 and 1.429802 gramme. The latter number divided by the former gives 15.96 as the specific gravity of oxygen. Stas, in recent experimental researches on combining-proportions, finds the atomic weight of oxygen to be not 16, but 15.96.

Exceptions to the law occur in a few compounds and in arsenicum and phosphorus, whose vapor-densities are twice that indicated by the rule. Possibly in these cases the temperature employed is insufficient to dissociate an unusually complex molecule into molecules of usual complexity. As regards compounds, and, possibly, as regards those elements in which the observed density is only half that indi-

cated by the rule, heat may, and in some cases probably does, produce molecular *dissociation* (*thermolysis*) into free atoms (uniatomic molecules) or into less complex molecules.

*Relation of the Specific Heat of Elements to their Atomic Weights.*—Reference may here appropriately be made to a physical fact of great importance as regards molecular and atomic weights. In the earlier pages of this manual it was stated that elements do not combine chemically in haphazard proportions, but in fixed weights; and abundant evidence of the truth of the statement has already been afforded, and will also be found in this section on quantitative analysis. Secondly, it has been shown that elements do not combine in haphazard proportions by volume, but in certain constant bulks; and the weights of these bulks have been found to be identical with the combining weights themselves. Thirdly (this is the point to which attention is now drawn), if equal amounts of heat be given to elements in the solid state (that is, to solid elements or to solid compounds of volatile elements), and the quantity of the element be increased or diminished until each is thus heated through an equal number of degrees, it will be found that the different weights of elements required are (in relation to a common standard) identical with the combining weights of the elements, and with the weights of the combining volume of the elements. Thus, where 108 parts of silver would be employed, 207 of lead would be necessary.* Hence, in the determination of (*a*) combining proportion, (*b*) specific gravity in gaseous state, and (*c*) specific heat, three distinct methods of ascertaining atomic weight are available. In cases where one method is inapplicable, recourse is had to either or, if practicable, both of the others, and thus the trustworthiness of observations and generalizations placed more or less beyond question. The specific heat of a solid element is the same in the free as in the combined condition; therefore the specific heat of a molecule is the sum of the specific heat of its constituent atoms. From the specific heat of a solid compound of a volatile element (chlorine, for example) can thus be calculated the specific heat of an element in the solid state, even though the free element cannot itself be solidified. For the processes by which experimentally to determine specific heat the reader is referred to books on physics.

There is equivalency, also, between electrical and chemical action. The amount of electricity which would set free 127 parts of iodine would set free 80 parts of bromine.

---

* Obviously, if equal weights of silver and lead were heated through an equal number of degrees, the silver would absorb nearly twice as much heat as the lead. In fact, as regards all the solid elements, "specific heats and atomic weights are inversely proportional." This law was discovered by Dulong and Petit. It follows that the product of the multiplication of the figures representing the specific heat of an element with the figures representing its atomic weight is, in the case of every element given, the same number.

## QUESTIONS AND EXERCISES.

971. Define specific weight, or, as it is commonly termed, specific gravity.

972. In speaking of light and heavy bodies especially, what standard of comparison is conventionally employed?

973. How are specific gravities expressed in figures?

974. Why should specific gravities be taken at one constant temperature?

975. How does the buoyancy of air affect the real weight of any material?

976. Describe the difference between density and specific gravity.

977. Give a direct method for the determination of the specific gravity of liquids.

978. A certain bottle holds 150 parts, by weight, of water, or 135.7 of spirit of wine; what is the specific gravity of the latter? *Ans.* 0.9046.

978*a*. An imperial fluidounce of a liquid weighs 366½ grains; what is its specific gravity? *Ans.* .838.

979. Equal volumes of bonzol and glycerine weigh 34 and 49 parts respectively, and the sp. gr. of the benzol is 0.850; what is the specific gravity of the glycerine? *Ans.* 1.225.

980. Explain the process employed in taking the specific gravity of solid substances in mass and in powder.

981. State the method by which the specific gravity of a light body, such as cork, is obtained.

982. What modifications of the usual method are necessary in ascertaining the specific gravity of substances soluble in water?

983. How is the specific gravity of gases determined?

984. By what law can the volume of a gas, at any required pressure, be deduced from its observed volume at another pressure?

985. To what extent will 78 volumes of a gas at 29.3 inches barometer alter in bulk when the pressure, as indicated by the barometer, is 30.2 inches?

986. Write a short account of the means by which the volumes of gases are corrected for temperature.

987. At the temperature of 15° C. 40 volumes (litres, pints, ounces, cubic feet, or other quantity) of a gas are measured. To what extent will this amount of gas contract on being cooled to the freezing-point of water (0° C.)?

*Answer.* As 1 vol. of any gas *at zero* expands or contracts .003665 of a vol. for each rise or fall of 1° C., 1 vol. at 0° C. if heated to 15° C. will become increased by 054975 (that is .003665 multiplied by 15), 1 vol. will expand to 1.054975. Conversely, 1 054975 vol. will contract to 1 vol. if cooled from 15° C. to 0° C. And if 1.054975 becomes 1 in cooling through 15° C., 40 vols. will (as found by rule-of-three) contract to 37.916.

(The following five problems and solutions are from Williamson's "Chemistry.")

988. 10 litres of oxygen are measured off at 14° F. Required the volume of the gas at 15° C.

*Answer.* The first operation must be to reduce the temperature quoted in Fahrenheit's degrees to an equivalent value on the Centigrade scale. 14° F. is 18° below 32° F., the freezing-point of water: and a range of 9° on the Fahrenheit's scale is equal to a range of 5° on the Centigrade scale, so that the temperature at which the oxygen is measured off is —10° C. The rise of temperature up to 0° expands the gas in such proportion that its volume at 0° is to its volume at —10 as 1 is to 1 —0.03665, *i. e.* as 1 to 0.96335. The further rise of temperature from 0° C. to 15° expands the gas in the proportion of 1 to $1 + 15 \times 0.003665$; *i. e.* 1 to 1.054975. The total rise of temperature therefore expands the gas in the proportion of 0.96335 to 1.054975.

$$0.96335 : 1.054975 : : 10 : x;$$

$$\therefore x = \frac{10 \times 1.054975}{0.96335} = 10.95$$

989. 230 cubic centimetres of oxygen are measured off at 14° C. and 740 millimetres mercurial pressure. Required the volume of the gas at the normal temperature and pressure (0° C. and 760 millimetres).

*Answer.* Let the reduction for change of temperature be made first. The proportion

$$1 + 14 \times 0.003665 : 1 : : 230 : x$$

gives

$$x = \frac{230}{1.05131} = 218.774.$$

To reduce this volume of 740 millimetres pressure to the volume corresponding to the pressure of 760 millimetres, we have the proportion

$$38 : 37 : : 218.77 : x;$$

whence

$$x = \frac{37 \times 218.77}{38} = 213.02.$$

990. A litre of oxygen is confined in a glass flask at 10° C. by the atmospheric pressure, added to that of a column of mercury 60 millimetres high. The flask must be heated to 300° C. without any increase of volume taking place in the oxygen. How high must the column of mercury then be which presses on the gas, supposing the atmospheric pressure to remain constant at 760 millimetres?

*Answer.* The oxygen is given at 10° C. and 820 millimetres pressure. If the pressure remained constant, the rise of temperature from 10° C. to 300° C. would expand the gas in such proportion that 1.03665 volume would expand to 2.0995 volumes. In order to prevent any expansion the pressure must be increased in the same proportion, whence

$$1.03665 : 2.0995 : : 820 : x;$$

$$\therefore x = \frac{820 \times 2.0995}{1.03665} = 1660.6.$$

From this total pressure the atmospheric pressure of 760 millimetres has to be deducted, leaving 900.6 millimetres as the height of the required mercurial column.

991. A litre of oxygen is required of the density of 100 at 0° C. What weight of potassic chlorate must be used for its preparation, and what total pressure must be applied to it?

*Answer.* The pressure required to compress oxygen from the density of 16 to that of 100 is found by the proportion

$$16 : 100 : : 760 : x;$$

$$\therefore x = \frac{76000}{16} = 4750.$$

At the pressure of 4750 millimetres of mercury the weight of a litre of oxygen (16 grammes measure 11.2 litres at 0° C. and 760 millims. pressure) is found by the proportion

$$760 : 4750 : : \frac{16}{11.2} : x;$$

whence

$$x = \frac{16 \times 4750}{11.2 \times 760} = 8.93 \text{ grammes.}$$

The weight of chlorate required for the evolution of 8.93 grammes of oxygen is found from the proportion

$$48 : 122.5 : : 8.93 : x;$$

$$\therefore x = 22.8 \text{ grammes.}$$

992. What is the volume of 12 grammes of hydrogen at 15° C.?

*Answer.* One gramme of hydrogen measures 11.2 litres at 0° C., therefore 12 grammes measure $12 \times 11.2 = 134.4$ litres at 0°. To find their volume at 15° C. we have the proportion

$$1 : 1 + 15 \times 0.003665 : : 134.4 : x;$$

whence

$$x = 134.4 \times 1.054975 = 141.788 \text{ litres.}$$

993. What interest for chemists have the specific heats of substances?

---

## Quantitative Analysis.

*Preliminary Note.*—Great care should be observed in selecting a *fair* sample of any bulk of material that is to be examined either by volumetric or gravimetric quantitative analysis. If the whole quantity is in separate parcels, and there is any ground for believing that the parcels differ in quality, they should, if practicable, be carefully mixed, or, technically, "bulked." Small portions should be taken from different parts of the resulting heap and well mixed in a mortar or other vessel, or, in certain cases, dissolved, and the solution well stirred or shaken A specimen of the powder, or a portion of the solution, may then be selected for analysis.

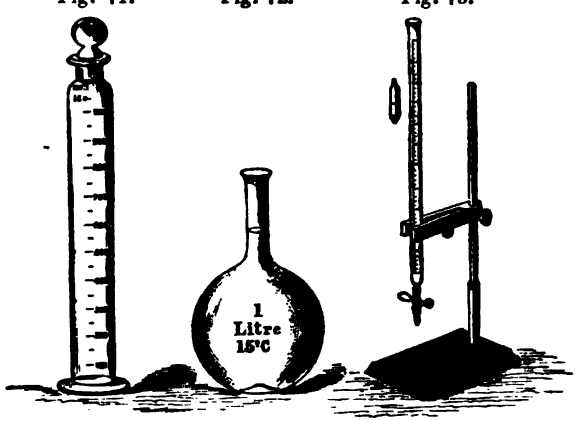
# VOLUMETRIC QUANTITATIVE ANALYSIS.

*(See Introductory Remarks to Quantitative Analysis.)*

## APPARATUS.

The only special vessels necessary in volumetric quantitative operations are: 1. A *litre flask* (Fig. 72), which, when filled to a mark on the neck, contains at 15° C., or about 60° F., one litre (1000 cubic centimetres, *i. e.* 1000 grammes of water*); it serves for preparing solutions in quantities of one litre. 2. A tall cylindrical *graduated litre jar* (Fig. 71) divided into 100 equal parts; it serves for the measurement and admixture of decimal or centesimal parts of a litre. 3. A graduated tube or *burette* (Fig. 73), which, when filled to 0, holds 100 cubic centimetres (a decilitre), and is divided into 100 equal parts; it is used for accurately measuring small volumes of liquids.

Fig. 71. Fig. 72. Fig. 73.




A litre jar. A litre flask. A burette, etc.

The best form of burette is Mohr's. It consists of a glass tube commonly about the width of a little finger and the length of an

* A cubic centimetre is, strictly speaking, the volume occupied by one gramme of distilled water at its point of greatest density, namely, 4° C.; metrical measurements, however, are uniformly taken at 15°.55 C. (60° F.).

arm from the elbow, contracted at the lower extremity and graduated. The width and length of burettes, however, as well as the extent and fineness of their graduation, vary considerably. To the contracted portion is fitted a small piece of vulcanized caoutchouc tubing, into the other end of which a small spout made of narrow glass tube is tightly inserted. A strong wire clamp effectually prevents any liquid from passing out of the burette unless the knobs of the clamp are pressed by the finger and thumb of the operator, when a stream or drops flow at will. In place of the India-rubber tubing and clamp, a stopcock is sometimes employed; and other modes of arresting the flow of liquid may be adopted. The accurate reading of the height of a solution in the burette is a matter of great importance; it should be taken from the bottom of the curved surface of the liquid. It may be still more exactly measured by the employment of a hollow glass float or bulb (Erdmann's float—see Fig. 73), of such a width that it can move freely in the tube without undue friction, and so adjusted in weight that it shall sink to more than half its length in any ordinary liquid. A fine line is scratched round the centre of the float; this line must always be regarded as marking the height of the fluid in the burette. In charging the burette, a solution is poured in, not until its surface is coincident with 0, but until the mark on the float is coincident with 0.

---

## ESTIMATION OF ALKALIES, ETC.

An equation represents much more than the formation of certain substances from others. Thus

$$\underset{\text{Ammonia.}}{2NH_4HO} + \underset{\substack{\text{Crystallized}\\\text{oxalic acid.}}}{H_2C_2O_4,2H_2O} = \underset{\substack{\text{Oxalate of}\\\text{ammonium.}}}{(NH_4)_2C_2O_4,H_2O} + \underset{\text{Water.}}{3H_2O}$$

not only shows that oxalate of ammonium is produced when ammonia and crystallized oxalic acid are mixed together, but, among other facts, that 70 parts of ammonia and 126 of oxalic acid yield 142 of crystallized oxalate of ammonium and 54 of water. For formulæ represent molecules; the weight of a molecule is the sum of the weight of atoms; and atomic weights are represented by definite invariable numbers (see the Table of atomic weights in the Appendix).

As 126 parts (= 1 molecule) of oxalic acid combined with 70 parts (= 2 molecules) of ammonia, 63 (half of 126) of oxalic acid will unite with 35 (= 1 molecule) of ammonia ($NH_4HO$); 63 parts of oxalic acid will also unite with 56 of caustic potash ($KHO = 56$), 40 of caustic soda ($NaHO = 40$), 100 of acid carbonate of potassium ($KHCO_3 = 100$), 69 of anhydrous carbonate of potassium ($K_2CO_3 = 138$), 84 of acid carbonate of sodium ($NaHCO_3 = 84$), 53 of anhydrous carbonate of sodium ($Na_2CO_3 = 106$), or 143 of crystallized carbonate of sodium ($Na_2CO_3,10H_2O = 286$). And if 63 parts of pure oxalic acid be dissolved in 100 volumes of water, the stated weights of these various salts should be exactly neutralized by such

a "standard" solution. 143 parts of crystallized carbonate of sodium, for instance, should, if pure, be exactly neutralized by the 100 volumes of the oxalic acid solution; and if a less number of volumes is required, the salt is so much per cent. impure. 143 parts by weight of a commercial sample of carbonate of sodium (common washing "soda") requiring only 97 of the standard oxalic acid solution is thus shown to contain 97 per cent. of pure carbonate of sodium, the remainder being impurities. Further, the strength of solutions of ammonia, soda, potash, and lime may be accurately determined by adding to any definite quantity of them gradually, from a burette, a solution containing oxalic acid in known quantity, until exact neutralization is effected. If the quantity of oxalic acid required be 63 parts by weight (or 100 volumes of solution of oxalic acid containing 63 parts by weight, of the crystals), then the quantities of alkaline solutions employed contain, of potash (KHO) 56 parts by weight, of soda (NaHO) 40, of ammonia ($NH_4HO$) 35 parts, of slaked lime, (Ca2HO) 37, anhydrous lime (CaO) 28, etc. The strength of the alkaline solution, or, in other words, the proportion required to effect neutralization of 100 volumes of the oxalic acid solution, having once been determined and decided by authority (B. P. *e. g.*), that quantity may be expected to take 100 volumes of the oxalic acid liquid; if less is required, the alkaline liquid is so much per cent. weaker than such official liquor.

In the following paragraphs, relating to the quantitative analysis of various alkaline substances by aid of a standard solution of oxalic acid, the acid is assumed to be quite pure. But oxalic acid of such *chemical* purity is very difficult to obtain. This difficulty may be got over by taking care that 100 volumes of the oxalic acid solution contain, by experiment, enough acid to exactly neutralize 53 parts of pure anhydrous carbonate of sodium. The latter is *not* difficult to obtain (see page 551). The oxalic solution is made a little stronger than necessary, then "set" with the carbonate, and afterwards diluted to whatever extent may be indicated by the experiment.

Another difficulty attends the use of the oxalic acid. It is liable to decompose if kept for any great length of time, especially if exposed much to light. This difficulty is only got over by substituting standard sulphuric acid, which is "set" or "standardized" in a similar manner.

In the experiments alluded to in the following sections, the volumetric experiments are made on rather a large scale, as recommended in the British Pharmacopœia, doubtless for use in drug laboratories. In such cases the experimental error would be slight, even if a standard solution were not absolutely chemically pure. Operations on the smaller scale of the chemical laboratory necessitate more delicate work, hence the recommendations to use sulphuric instead of oxalic acid, and to "set" with carbonate of sodium.

The exact point of neutralization of acid or alkaline liquids is experimentally ascertained by litmus paper, or, more generally, infusion of litmus, which is turned red by the slightest amount of free acid, and blue by alkali.

## Standard Solution of Oxalic Acid.

### (Crystallized Oxalic Acid, $H_2C_2O_4,2H_2O = 126$.)

On account of the bivalent character of the oxalic radical and the univalent character of most of the metals contained in the salts which are estimated by oxalic acid, it is convenient to take half the molecular weight of the acid for experiments, with the whole of the molecular weights of salts of univalent basylous radicals. The oxalic acid must be pure, leaving no ash when a gramme or so is heated to redness in a porcelain or platinum crucible; it must also be quite dry, but not effloresced.

Place 63 grammes of the crystals in a litre flask, add distilled water and shake till dissolved, diluting until the solution, at about 60° F., has an exact volume of 1 litre. Preserve in a stoppered bottle.

100 cubic centimetres of this solution contain $\frac{1}{10}$ of the molecular weight of oxalic acid in grammes, and will neutrallze $\frac{1}{10}$ of the molecular weight in grammes, of a salt whose molecule contains one atom of certain bivalent metals (as Ca2HO), or a salt ($Na_2CO_3$ *e. g.*) containing in one molecule two atoms of univalent metals, or $\frac{1}{10}$ of the molecular weight in grammes of salts, single molecules of which contain one atom of univalent radicals (such as $KHCO_3$).

The following official substances are tested with this solution. In those which are fluid there is commonly a slight variation in strength according as they are made by formulæ of the British or U. S. Pharmacopœia.

| | Grammes of substance. | C. c. of vol. sol. |
|---|---|---|
| *Ammonia, solution of . . . . . | 17.00 = | 100.0 |
| " strong solution of . . . . . | 5.23 = | 100.0 |
| Ammonium, carbonate of, B. P. and U. S. P. | 5.90 = | 100.0 |
| Borax, B. P. and U. S. P. . . . . | 19.10 = | 100.0 |
| Lead, acetate of, B. P. and U. S. P. . . . | 9.50 = | 50.0 |
| " sol. of subacetate of, B. P. and U. S. P. | 51.02 = | 100.0 |
| Lime, aqueous solution of, B. P. and U. S. P. | 438.00 = | 20.0 |
| " saccharated sol. of . . . . . | 45.30 = | 25.0 |
| Potash, caustic, B. P. and U. S. P. . . . | 5.60 = | 90.0 to 100.0 |
| " solution of . . . . . | 48.02 = | 50.0 |
| " water, effervescing . . . . . | 292.00 = | 10.0 |

* The reading of the Table may be amplified thus: 17 grammes of the official solution of Ammonia (*Liquor Ammoniæ*, B. P.), carefully weighed, will require, for complete neutralization, if of full strength, 100 cubic centimetres of the official volumetric solution of oxalic acid.

| | Grammes of substance. | C. c. of vol. sol. |
|---|---|---|
| Potassium, bicarbonate of, B. P. and U. S. P. | 5.00 = | 50.0 |
| " acid tart. of, B. P. and U. S. P. | 18.80 = | 100.0 |
| " carbonate of, B. P. and U. S. P. | 8.30 = | 98.0 to 100.0 |
| " citrate of, B. P. and U. S. P. . | 10.20 = | 100.0 |
| " tartrate of, B. P. and U. S. P. | 11.30 = | 100.0 |
| Soda, caustic, B. P. and U. S. P. . . . | 4.00 = | 90.0 to 100.0 |
| " solution of . . . . . | 48.72 = | 50.0 |
| " water, effervescing . . . . . | 246 07 | 10.0 |
| Sodium, bicarbonate of, B. P. and U. S. P. | 8.40 = | 100.0 |
| " and potassium, tart. of, B. P. and U. S. P. | 14.10 = | 100.0 |
| " carbonate of, B. P. and U. S. P. . | 14.30 = | 96.0 to 100.0 |

*Note* 1.—The several substances diluted or dissolved, as described in the following paragraphs, are conveniently placed in a beaker, and the solution of acid run in cautiously from the burette.

*Note* 2.—A smaller number of c. c. of volumetric solution than 100, and a corresponding amount of substance to be tested, may be employed in any case. But to ascertain percentage of impurity, the amount of substance corresponding to 100 c. c. of the vol. sol., and a porportionate amount of the vol. sol. itself, must be considered to have been used, the number of c. c. then wanting to make up 100 is the percentage of impurity. Here and elsewhere in quantitative analysis, facility in the use of the "rule of three" saves much time and trouble.

*The solutions of ammonia* require only the addition of solution of litmus, and the acid cautiously added until the last drop turns the liquid red. The amount of acid used previously to the addition of the portion that reddened the litmus indicates the proportional purity of the alkaline liquid. Thus, if only 50 c. c. are required, the solution is only half as strong as it ought to be; if 93 c. c. are needed, the sol. of ammonia is 7 per cent. too weak, and so on. The actual quantity of ammonia ($NH_4HO$) or ammoniacal gas ($NH_3$) in the solutions is readily ascertained by calculations, thus: 100 c. c. of the acid solution have been employed; these contain $\frac{1}{2}\frac{1}{0}$ of the molecular weight of oxalic acid in grammes = 6.3, and have neutralized $\frac{1}{1}\frac{1}{0}$ of the molecular weight of ammonia in grammes = 3.5 (or 1.7 of $NH_3$); 5.23 parts, by weight, of strong solution of ammonia (the amount employed in the experiment) contain, therefore, 3.5 of ammonia, $NH_4HO$, or 1.7 of ammoniacal gas, $NH_3$; now if 5.23 parts of a solution contain 3.5 of real ammonia ($NH_4HO$), 100 parts will contain (by rule-of-three calculation) 67, and if the 5.23 contain 1.7 parts of ammonia gas ($NH_3$), 100 will contain 32.5; hence the Strong Solution of Ammonia, supposed to have been under examination, contains 67 per cent. of the hydrate or 32.5 per cent. of the

gas.* The formulæ and molecular weights representing this process are as follows :—

$$\underbrace{2NH_4HO}_{70} \quad \underbrace{(\text{or } 2NH_3)}_{34} \quad + \quad \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*The carbonate of ammonium* should be dissolved in 30 or 40 c. c. of distilled water, infusion of litmus added, the standard oxalic acid solution allowed to flow in until the well-stirred liquid assumes a purplish hue (due to the influence of carbonic acid on the litmus), the whole gently warmed to promote evolution of carbonic acid gas, more standard acid then dropped in until the liquid again becomes purple, heat once more applied, and the operation continued until the last drop of acid turns the solution red; the height of the column of liquid in the burette before the last drop escapes represents the true amount of standard solution used, and hence the percentage of real carbonate of ammonium (of official quality) in the specimen on which the experiment was performed. The solution must be heated with care, or ammonia will escape. The commercial carbonate of ammonium now commonly met with, is more nearly a mixture of one molecule of acid carbonate to one of carbamate of ammonium (Divers) rather than two of the former to one of the latter, as shown in the British Pharmacopœia; 5.9 grammes would, therefore, require more nearly 112 or 113 c. c. rather than 100 c. c. of the standard solution of oxalic acid. (Practised analysts usually add excess of the standard acid and thus fix every trace of ammonia; then gently boil to

---

* Extremely minute quantities of ammonia—1 part in many millions of water—may be estimated volumetrically by adding excess of a colorless solution of red iodide of mercury (Nessler's test), then in a similar vessel, containing an equal amount of pure water with excess of the Nessler reagent, imitating the depth of yellow or reddish-yellow color thus produced by adding an ammoniacal solution *of known strength*. The amount of ammonia thus added represents the amount in the original liquid.

*The Nessler Reagent.*—A litre may be made by dissolving 30 or 40 grammes of iodide of potassium in a small quantity of hot water, adding a strong hot solution of perchloride of mercury until the precipitate of mercuric iodide ceases to redissolve by the aid of rapid stirring and heat, slightly diluting, filtering, adding a strong solution of (120 to 140 grammes of) caustic soda or (160 to 180 grammes of) caustic potash, and diluting to 1 litre. A few c. c. (5 or 6 or more) of a strong solution of perchloride of mercury are finally stirred in, the whole set aside till all precipitated red iodide has deposited, and the clear liquid decanted for use. The reaction of this Nessler test with ammonia is as follows :—

$$NH_3 + 2HgI_2 + 3KHO = NHg_2I + 3KI + 3H_2O.$$

get rid of carbonic acid gas; bring back the liquid to neutrality by an observed volume of standard alkaline solution, and deduct an equivalent volume of acid from the quantity first added.) The formulæ and weights representing the process are as follows:—

$$\underbrace{N_4H_{16}C_8O_8}_{236} + \underbrace{2(H_2C_2O_4,2H_2O)}_{252}$$

(*vide* p. 92) (2 molecules)

$$\underbrace{\text{or } 2N_3H_{11}C_2O_5}_{314} + \underbrace{3(H_2C_2O_4,2H_2O)}_{378}$$

$$236 \ldots\ldots\ldots 248$$

$$5.9 \ldots\ldots\ldots 7.1$$

As 252 parts of oxalic acid neutralize 236 of the so-called carbonate of ammonium, 6.3 of acid (or 100 vols. of its solution) will neutralize 5.9 of carbonate. If 5.9 grammes of carbonate be the quantity employed, and it does *not* require 100 c. c. of the volumetric solution, it is so much per cent. weak. Thus, if say 94 c. c. neutralize the salt, the latter is 6 per cent. weak (some ammonia gas has escaped, and consequently, an abnormal amount of acid carbonate $NH_4HCO_3$, is present). Any smaller quantity of salt than 5.9 grammes may be used; in that case a rule-of-three sum must be worked to show how many c. c. of vol. sol. would have been employed if 5.9 grammes had actually been the amount under experiment.

*The borax* should be dissolved in several ounces of distilled water. The formulæ and molecular weights representing the process are as follows:—

$$\underbrace{2NaBO_2,B_2O_3,10H_2O}_{382} + \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*The solutions of the acetates of lead* in distilled water may be rendered clear by the addition of a few drops of acetic acid. They must be well stirred after each addition of the solution of oxalic acid. The action is complete when the last drop of acid produces no more precipitate (oxalate of lead).

$$\underbrace{Pb2C_2H_3O_2,3H_2O}_{379} + \underbrace{H_2C_2O_4,2H_2O}_{126}$$

$$\underbrace{Pb_2O2C_2H_3O_2}_{548} + \underbrace{2(H_2C_2O_4,2H_2O)}_{252}$$

*The solutions of lime* require similar treatment.

$$\underbrace{Ca2HO}_{74} \quad \underbrace{(\text{or } CaO)}_{56} \quad + \quad \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*Solid caustic potash or soda* is never met with in a state of chemical purity, but should contain not less than 90 per cent. of the hydrate of potassium or sodium. The standard acid is added to an aqueous solution of the hydrate, the termination of the action between the alkali and acid being observed by aid of litmus.

If carbonic acid be present, the mixture must be gently boiled before a final reading of the amount of acid added is taken.

$$\underbrace{2KHO}_{112} \quad \text{or} \quad \underbrace{2NaHO}_{80} \quad + \quad \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*The alkaline carbonates* are often moist, and include traces of sulphates, chlorides, and silicates, but are sufficiently pure if containing, in the case of carbonate of potassium 98 per cent., and carbonate of sodium 96 per cent., of the respective crystalline salts. The volumetric manipulations with these salts are similar to those for carbonate of ammonium. The strength of soda-ash is often reported in terms of "soda," that is, oxide of sodium ($Na_2O$=62). The old molecular weight of carbonate of sodium, 54 (it should have been 53), derived from that of "soda," 32 (it should have been 31), is still employed in Great Britain in reporting the strength of soda-ash. The true amount of soda equivalent to 54 parts of carbonate is 31.41 parts. A modern analyst, having found the true amount of soda in a sample of soda-ash, is expected by some manufactures to report 31.41 parts for every 31, and 54 of carbonate instead of 53, and other quantities in proportion to these figures.

$$\underbrace{2KHCO_3}_{200} + \underbrace{H_2C_2O_4,2H_2O}_{126} \qquad \underbrace{2NaHCO_3}_{168} + \underbrace{H_2C_2O_4,2H_2O}_{126}$$

$$\underbrace{K_2CO_3}_{138} + \underbrace{H_2C_2O_4,2H_2O}_{126} \qquad \underbrace{Na_2CO_3}_{106} + \underbrace{H_2C_2O_4,2H_2O}_{126}$$

$$\underbrace{K_2CO_3 \text{ of } 14\ \%}_{164.285} + \underbrace{H_2C_2O_4,2H_2O}_{126} \qquad \underbrace{Na_2CO_3,10H_2O}_{286} + \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*One molecule of tartrate of potassium*, or two of the *acid tartrate*, yields one of carbonate when burnt, two molecules of

*citrate*, yielding three of carbonate under the same circumstances. Two molecules of *tartrate of potassium and sodium* yield one of potassium carbonate and one of sodium carbonate. A volumetric estimation of the amount of carbonate thus produced affords indirect means of quantitatively determining the purity of the original salts.

$$\underbrace{K_2C_4H_4O_6}_{226} \text{ eq. to } \underbrace{H_2C_2O_4,2H_2O}_{126}$$

$$\underbrace{2KHC_4H_4O_6}_{376} \text{ eq. to } \underbrace{H_2C_2O_4,2H_2O}_{126}$$

$$\underbrace{2K_2C_6H_5O_7}_{612} \text{ eq. to } \underbrace{3H_2C_2O_4,2H_2O}_{378}$$

$$\underbrace{NaKC_4H_4O_6,4H_2O}_{282} \text{ eq. to } \underbrace{H_2C_2O_4,2H_2O}_{126}$$

*Alkalimetry.*—The foregoing processes are often spoken of as those of *alkalimetry* (the measurement of alkalies).

---

*Notes.*

*Neutral solution of litmus* is prepared by digesting the commercial fragments in about fifteen or twenty times their weight of water for a few hours, decanting, dividing into two equal portions, adding acid to one till it is faintly red, then pouring in the other, and mixing. The solution may be kept in a stoppered bottle, but occasionally exposed to the air. It should never be filtered, but gradually allowed to deposit.

*Standard sulphuric acid* may be used in the place of oxalic acid if the latter cannot readily be obtained in a state of purity, 100 c. c. of the liquid containing $\frac{1}{20}$ of the molecular weight of the pure acid in grammes. It is prepared by diluting oil of vitriol with from three to four times its bulk of distilled water, ascertaining how much of the acid liquid is required to exactly neutralize $\frac{1}{20}$ of the molecular weight of pure carbonate of sodium, taken in grammes (5.3), and adding water until the observed volume of acid is increased to 100 c. c. Chemically pure anhydrous carbonate of sodium ($Na_2CO_3$) is obtained by heating the pure bicarbonate to dull redness in a platinum or porcelain crucible for about a quarter of an hour. The commercial bicarbonate should be tested for chlorides and sulphates,

which are usually present in small quantities. Two or three hundred grammes may be purified by washing first with a saturated solution of bicarbonate of sodium and then cold distilled water until all trace of impurity has disappeared, drying over a water-bath, and then igniting to convert into carbonate. Chemically pure carbonate of potassium and bicarbonate of potassium may be obtained in a similar manner.

*Other quantities of salts* than those stated in the foregoing and following Tables may be employed in volumetric determinations, calculation giving any desired form to the experimental results, an expert analyst thus saving much time and material. In the case of substances which are liable to alter by exposure to air, it is important that a selected quantity should be quickly weighed, rather than selected weights be accurately balanced by material, the former operation occupying much the shorter time.

*Salts other than the official* may be quantitatively analyzed by the volumetric solutions, slight modifications of manipulation even enabling the processes to be adapted to fresh classes of salts. Ample instructions for extending operations in this manner will be found in Sutton's "Handbook of Volumetric Analysis."

## QUESTIONS AND EXERCISES.

994. Describe the various pieces of apparatus used in volumetric determinations.

995. One hundred cubic centimetres of solution of oxalic acid contain 6.3 grammes of the crystallized salt; what weights of bicarbonate of potassium and anhydrous carbonate of sodium will that volume saturate?—*Ans.* 10 grammes and 5.3 grammes.

996. What weight of hydrate of potassium is contained in solution of potash 48.02 grammes of which are saturated by 50 c. c. of the standard solution of oxalic acid?—*Ans.* 5.83 per cent.

997. State the percentage of hydrate of calcium in lime-water 438 grammes of which are neutralized by 20 c. c. of the volumetric solution of oxalic acid.—*Ans.* .1689.

998. Eight grammes of a sample of Rochelle salt, after ignition, etc., require 54.3 c. c. of the official oxalic acid solution for complete saturation; what is the centesimal proportion of real salt present?—*Ans.* 95.7.

## ESTIMATION OF ACIDS.

In the previous experiments a known amount of an acid has been used in determining unknown amounts of alkalies. In those about to be described a known amount of an alkali is employed in estimating unknown amounts of acids. The alkaline salt selected may be

either a hydrate or a carbonate; but the former is to be preferred; for the carbonic acid, set free when a strong acid is added to a carbonate, interferes to some extent with the indications of alkalinity, acidity, or neutrality afforded by litmus. The alkali most convenient for use is soda, a solution of which has probably already been made the subject of experiment in operations with the standard solution of oxalic acid. It should be kept in a stoppered bottle, and exposed to air as little as possible.

### Standard Solution of Soda.

(Hydrate of Sodium, NaHO = 40.)

100 c. c. of the standard solution of oxalic acid are placed in a beaker with a little litmus, the tube or burette in which the acid was measured rinsed out, and the washings poured into the beaker. A little strong solution of caustic soda is poured through the burette to rinse out water adhering to the tube and float (these rinsings thrown away), and the tube then filled to 0 with more of the alkaline liquid. The solution of soda is cautiously allowed to flow into the beaker until exact neutrality is obtained, the quantity noted, and, to every similar quantity of the whole bulk of the solution of soda, water added until the liquid measures 100 parts. If, for example, 93 c. c. of solution of soda have neutralized the 100 c. c. of acid, then 7 c. c. of distilled water must be added to 93 c. c. of the soda solution, or 70 to 930 to make a litre. A sum of simple proportion will show to what extent any other quantity is to be diluted. Thus, if the bulk of soda solution remaining measures, say, 900 c. c., its volume must be augmented to 967.7 c. c.; for if 93 are to be diluted to 100, 900 must be diluted to 967.7.

100 c. c. of the soda solution contain $\frac{1}{10}$ of the molecular weight (= 4), taken in grammes, of pure hydrate of sodium, and will neutralize an equivalent quantity of any acid. That is, 100 c. c. will neutralize $\frac{1}{10}$ of the molecular weight in grammes of an acid whose molecule contains one atom of any univalent acidulous radical, $\frac{1}{20}$ of the molecular weight in grammes of an acid a molecule of which contains one atom of any bivalent acidulous radical, or $\frac{1}{30}$ of the molecular weight in grammes of an acid single molecules of which contain one atom of any trivalent acidulous radical.

The following official acids are tested with this solution:—

| | Grammes of substance. | | C. c. of vol. sol. |
|---|---|---|---|
| Acid, acetic . . . . . | 18.20 | = | 100.0 |
| " " diluted, B. P. and U. S. P. | 70.29 | = | 50.0 |
| " " glacial . . . . . | 6.00 | = | 99.0 |
| " citric, B. P. and U. S. P. . . | 7.00 | = | 100.0 |
| " hydrochloric, B. P. and U. S. P. | 11.48 | = | 100.0 |
| " " diluted . . . . | 34.50 | = | 100.0 |
| " nitric, B. P. and U. S. P. . . | 9.00 | = | 100.0 |
| " " diluted . . . . . | 36.13 | = | 100.0 |
| " nitro-hydrochloric, diluted . . | 38.30 | = | 100.0 |
| " sulphuric, B. B. and U. S. P. . | 5.06 | = | 100.0 |
| " " aromatic . . . . | 36.35 | = | 100.0 |
| " " diluted . . . . . | 35.90 | = | 100.0 |
| " tartaric, B. P. and U. S. P. . . | 6.50 | = | 100.0 |

*Notes.*—1. In volumetrically estimating the strength of acids by an alkali, the indicator of neutrality is the same as that used in testing alkalies by an acid, namely, litmus.

2. Pure acetates, citrates, tartrates, and some other organic salts have an alkaline action on litmus, but not to an important extent. If the soda solution be added to acetic, citric, or tartaric acids, containing litmus, until the liquid is fairly blue, the operator will obtain trustworthy results. In delicate experiments turmeric may be used instead of litmus.

3. Six grammes of *pure* glacial acetic acid are neutralized by 100 c. c. of the standard solution of soda. But acid of this degree of purity is extremely difficult to prepare. The commercial acid contains only 1 per cent. of water, and is sufficiently pure for use in medicine.

*Acidimetry.*—The operations for the quantitative analysis or measurement of acids are often collectively spoken of under the name of *acidimetry*. They admit of considerable extension. (See the work by Sutton previously cited.)

*Percentage strength of Acids.*—The percentage strength of the several acids and their official solutions is readily ascertained by calculation in a manner similar to that given for alkalies. Thus the volumetric operation with the liquid commonly termed acetic acid is based on the reaction expressed in the following equation:—

$$\underset{\text{Acetic acid.}}{HC_2H_3O_2} + \underset{\text{Soda.}}{NaHO} = \underset{\text{Acetate of sodium.}}{NaC_2H_3O_2} + \underset{\text{Water.}}{H_2O.}$$

This equation, translated into parts by weight, means that 60 parts (the molecular weight) of true acetic acid are exactly neutralized by 40 parts (the molec. wt.) of soda. Supposing the quantity of "acetic acid" employed in the experiment to have been that recommended in the Table (18.2 grammes), and that to neutralize it 100 c. c. of the soda solution have been used, and remembering that 100 c. c. of the

soda solution contain 4 grammes of soda, it follows that the 18.2 grammes of the liquid called acetic acid contains 6 grammes of real acetic acid; for if 40 (the molec. wt.) of soda neutralize 60 (the molec. wt.) of real acetic acid, 4 will neutralize 6. Lastly, if 18.2 contain 6, 100 will contain 33; the so-called acetic acid (really an aqueous solution) contains 33 per cent. of true acetic acid ($HC_2H_3O_2$).

If the 18.2 grammes have taken, say, 93 instead of 100 c. c, the acid liquid is 7 per cent. weak; in other words, it contains only 30.7 per cent. of real acid; for (by rule-of-three) if 100 c. c. added to 18.2 grammes indicate 33 per cent., 93 c. c. added to 18.2 grammes indicate 30.7 per cent.

*The remaining acids* react with the alkaline salts in the manner and to the extent indicated by the following formulæ and molecular weights:—

$$\underbrace{H_2C_6H_6O_7,H_2O}_{210} + \underbrace{3NaHO}_{120}$$

$$\underbrace{HCl}_{36.5} + \underbrace{NaHO}_{40} \qquad \underbrace{HNO_3}_{63} + \underbrace{NaHO}_{40}$$

$$\underbrace{H_2SO_4}_{98} + \underbrace{2NaHO}_{80} \qquad \underbrace{H_2C_4H_4O_6}_{150} + \underbrace{2NaHO}_{80}$$

## QUESTIONS AND EXERCISES.

999. What percentage of real acid is present in diluted sulphuric acid 30 grammes of which are neutralized by 84 c. c. of the official volumetric solution of soda?—*Ans.* 13.72.

1000. How much real nitric acid is contained in a solution 36 grammes of which are saturated by 94 c. c. of the standard solution of soda?—*Ans.* 16.45 per cent.

## ESTIMATION OF ACIDULOUS RADICALS PRECIPITATED BY NITRATE OF SILVER.

The purity of many salts and the strength of their solutions may be determined by this process; but at present only three official substances (namely, diluted hydrocyanic acid, bromide of potassium, and arseniate of sodium) are quantitatively analyzed by standard solution of nitrate of silver. The reactions on which the success of the process depends are expressed in the following equations:—

$$AgNO_3 + 2NaCy = NaCyAgCy + NaNO_3,$$

then, $NaCyAgCy + AgNO_3 = 2AgCy + NaNO_3$.

$$KBr + AgNO_3 = AgBr + KNO_3.$$

$$Na_2HAsO_4 + 3AgNO_3 = Ag_3AsO_4 + 2NaNO_3 + HNO_3.$$

## Standard Solution of Nitrate of Silver.

(Nitrate of Silver, $AgNO_3 = 170$.)

Dissolve 17 grammes of crystals of pure nitrate of silver in 1 litre of water. 100 c. c. of this solution contain $\frac{1}{100}$ of the molecular weight in grammes of nitrate of silver, and will decompose an equivalent quantity of a salt of any acidulous radical yielding silver compounds insoluble in water.

| | Grammes of substance. | C. c. of vol. sol. |
|---|---|---|
| Acid, hydrocyanic, diluted, B. P. and U. S. P. . | 27.00 | = 100.0 |
| Potassium, bromide of, B. P. and U. S. P. . . | 1.19 | = 100.0 |
| Sodium, arseniate of (dry) . . . . . | .62 | = 100.0 |

*Diluted hydrocyanic acid* is converted into cyanide of sodium by adding caustic soda until, after stirring, litmus shows that the liquid has a decided but not an excessive alkaline reaction. The resulting fluid is well diluted. The nitrate-of-silver solution is then allowed to flow in gradually, until, after thorough agitation, a slight permanent turbidity obtains—the liquid remaining fairly but not excessively alkaline throughout and especially at the end of the operation, as shown by an immersed piece of litmus-paper.

Reference to the foregoing equations show that one molecular weight of nitrate of silver is used by *two* molecular weights of cyanide of sodium in the formation of the double compound. 100 c. c. of standard solution contain $\frac{1}{100}$ of the molecular weight in grammes of nitrate of silver (= 1.7), and correspond therefore to $\frac{1}{100}$ of *two* molecular weights in grammes of hydrocyanic acid (= 0.54).

$$\underbrace{2HNC}_{54} \text{ eq. to } \underbrace{AgNO_3}_{170}$$

*Bromide of potassium* is dissolved in distilled water in a beaker, and the standard solution added until, after agitation of the liquid and subsidence of the bromide of silver, a drop of the solution of nitrate of silver gives no more precipitate.

$$\underbrace{KBr}_{119} + \underbrace{AgNO_3}_{170}$$

*Arseniate of sodium* ($Na_2HAsO_4,7H_2O$) must be dried at 300° F. before weighing. It is thus reduced to a definite

anhydrous salt ($Na_2HAsO_4$), losing, if pure, 40.38 per cent. of water. Of the weighed residue .62 gramme is dissolved in distilled water, 3.3 c. c. of the vol. sol. of soda added, and the whole treated as described in the previous paragraph.

$$\underbrace{Na_2HAsO_4}_{186} + NaHO + \underbrace{3AgNO_3}_{510} = Ag_3AsO_4 + 3NaNO_3 + H_2O$$

*Spirit of Wine* (*Spiritus Rectificatus*, B. P.) may contain traces of amylic alcohol and aldehyd; these may be detected by nitrate of silver, which is reduced by them to the metallic state. Any quantity beyond a mere trace of such bodies renders spirit of wine too impure for use in medicine. "Four fluidounces with thirty grain-measures (about two cub. cent.) of the volumetric solution of nitrate of silver exposed for twenty-four hours to bright light, and then decanted from the black powder which was formed, undergoes no further change when again exposed to light with more of the test."

*The Volumetric Estimation of Iodide of Potassium.*—M. Personne has proposed a method based upon the following reaction: If solution of perchloride of mercury be added, a little at a time, to a solution of iodide of potassium, chloride of potassium is formed together with a soluble compound of iodide of potassium and periodide of mercury. But when one-half the iodide of potassium has been decomposed by the perchloride of mercury, the smallest further addition of that compound produces a persistent red precipitate. Thus, $4KI + HgCl = 2KCl + HgI_2, 2KI$. And when every molecule of the iodide has thus been attacked, the further addition of the perchloride gives the permanent red precipitate of periodide, thus $HgI_2 2KI + HgCl_2 = 2HgI_2 + 2KCl$.

To prepare a semi-decinormal (p. 566) test solution, 13.55 grammes of perchloride of mercury and 8 or 10 grammes of chloride of potassium or sodium are dissolved in 300 or 400 grammes of distilled water; afterwards sufficient water is added to bring the quantity up to 1000 c. c. In another vessel 33.20 grammes of the iodide of potassium to be tested are to be dissolved in sufficient distilled water to form a litre of solution. Ten c. c. of the solution of iodide of potassium are then placed in a glass vessel, and the mercurial solution is added drop by drop from a burette, graduated to tenths of a cubic centimetre, the mixture being continually shaken. The reaction is terminated when the mercurial solution produces a perceptibly red and persistent precipitate. It is only necessary then to read off from the burette the number of divisions employed to ascertain the richness of the iodide. (If pure, the 10 c. c. will require 10 c. c.)

| | | | |
|---|---|---|---|
| $HgCl_2 = 271$ | or 27.1 | or 13.55 | or .1355 in 10 c. c. |
| $4KI = 664$ | or 66.4 | or 33.20 | or .3320 in 10 c. c. |

M. Personne has ascertained that the presence of carbonate, chloride, or bromide of potassium, is no obstacle to this application of volumetric analysis.

## QUESTIONS AND EXERCISES.

1001. Explain the volumetric method of estimating the strength of aqueous solutions of hydrocyanic acid.

1002. How much nitrate of silver will indicate, by the official volumetric process, the presence of 1 part of real hydrocyanic acid? —*Ans.* 3.148 parts.

---

## ESTIMATION OF SUBSTANCES READILY OXIDIZED.

Any deoxidizer, that is, any substance which quickly absorbs a definite amount of oxygen or is susceptible of any equivalent action, may be quantitatively tested by ascertaining how much of an oxidizing agent of known power must be added to a given quantity before complete oxidation is effected. The oxidizing agents employed for this purpose in the British Pharmacopœia are iodine and the red chromate of potassium; permanganate of potassium is often used for the same purpose. Iodine acts indirectly, by taking hydrogen from water and liberating oxygen; the red chromate of potassium directly, by the facility with which it yields three-sevenths of its oxygen—as indicated by the equations and statements given on page 561; permanganate of potassium by affording five-eighths of its oxygen in presence of acid, $2K_2Mn_2O_8 + 6H_2SO_4 = 2K_2SO_4 + 4MnSO_4 + 6H_2O + 5O_2$.

### Standard Solution of Iodide.

(Iodine, I = 127.)

Prepare pure iodine by mixing the commercial article with about a fourth of its weight of iodide of potassium and subliming. Sublimation may be effected by gently warming the mixture in a beaker, the mouth of which is closed by a funnel; the iodine vapor condenses on the funnel; while fixed impurities are left behind, and any chlorine which the iodide may contain is absorbed by the iodide of potassium, an equivalent quantity of iodine being liberated. Small quantities may be similarly treated between two watch-glasses, placed edge to edge. Any trace of moisture in the resublimed iodine is removed by exposure for a few hours under a glass shade near a vessel containing oil of vitriol.

Place 12.7 grammes of pure iodine and about 18 grammes of pure iodide of potassium (an aqueous solution of which is the best solvent of iodine; the salt plays no other part in these operations) in a litre flask, add a small quantity of water, and

agitate until the iodine is dissolved; dilute to 1 litre. 100 c. c. of this solution contain $\frac{1}{200}$ of the molecular weight of free iodine in grammes, and, water being present, will cause the oxidation of $\frac{1}{200}$ of the molecular weight of sulphurous acid ($H_2SO_3$) in grammes (= .41), or $\frac{1}{200}$ (= .32) of sulphurous acid gas ($SO_2$), sulphuric acid being formed. 100 c. c. will oxidize $\frac{1}{200}$ of the molecular weight of arsenious acid ($H_3AsO_3$) in grammes, or $\frac{1}{400}$ of the molecular weight of common white arsenic ($As_2O_3$) in grammes (= .495), arsenic acid ($H_3AsO_4$) being produced. 100 c. c. will also attack $\frac{1}{100}$ of the molecular weight of hyposulphite of sodium. The reactions are expressed in the following equations:—

$$I_2 + H_2O + H_2SO_3 = 2HI + H_2SO_4.$$
$$I_2 + 2H_2O + SO_2 = 2HI + H_2SO_4.$$
$$I_2 + H_2O + H_3AsO_3 = 2HI + H_3AsO_4.$$
$$2I_2 + 5H_2O + As_2O_3 = 4HI + 2H_3AsO_4.$$
$$I_2 + 2(Na_2S_2O_3,5H_2O) = 2NaI + Na_2S_4O_6 + 10H_2O.$$

The following official substances are tested with the standard solution of iodine:—

| | Grammes of substance. | | C. c. of vol. sol. |
|---|---|---|---|
| Acid, solution of sulphurous . . . . . | 3.47 | = | 100 |
| Arsenic, in mass, B. P. and U. S. P. . . . | 0.495 | = | 100 |
| " in alkaline sol. (*Liq. Arsenicalis*) . | 54.64 | = | 100 |
| " in acid sol. (*Liq. Arsen. Hydrochl.*) | 54.64 | = | 100 |
| Sodium, hyposulphite of . . . . . | 2.48 | = | 100 |

*The solution of sulphurous acid* is diluted with three-fourths of a litre of cold water, and the iodine solution added until a slight permanent brown tint is produced, showing the presence of free iodine. A better indication of the termination of the action is afforded by mucilage of starch, which gives a blue color with the slightest trace of iodine. As already stated, 100 c. c. of this volumetric solution contain an amount of free iodine sufficient to cause the oxidation of $\frac{1}{200}$ of the molecular weight of either sulphurous acid or sulphurous acid gas. Now $\frac{1}{200}$ of the molecular weight of sulphurous acid ($H_2SO_3$) in grammes is 0.41; if 3.47 grammes of the solution contain 0.41 of the acid, 100 of the solution will be found to contain 11.8. By a similar calculation the official solution may be shown to contain,.or, rather, yield 9.22 per cent. of sulphurous acid gas ($SO_2$).

If the sulphurous acid be diluted to a less degree than .04 or .05 per cent., there will be some risk of the sulphuric acid

formed being again reduced to sulphurous acid, with liberation of iodine. In delicate experiments the distilled water used for dilution should previously be freed from air by boiling, to prevent the small amount of oxidizing action which dissolved air would exert.

| $\underbrace{H_2SO_3}$ eq. to $\underbrace{I_2}$ | | $\underbrace{SO_2}$ eq. to $\underbrace{I_2}$ | |
|---|---|---|---|
| 82 | 254 | 64 | 254 |

*The solid arsenic* is dissolved in boiling water by help of about two grammes of bicarbonate of sodium. When the liquid is quite cold, mucilage of starch is added, and the iodine solution allowed to flow in until, after well stirring, a permanent blue color is produced.

| $\underbrace{H_3AsO_3}$ eq. to $\underbrace{I_2}$ | | $\underbrace{As_2O_3}$ eq. to $\underbrace{2I_2}$ | |
|---|---|---|---|
| 126 | 254 | 198 | 508 |

*The arsenical solution already containing some carbonate of potassium* requires only about one and a half gramme of bicarbonate of sodium for neutralization of the arsenious acid. After boiling and cooling, starch and the iodine solution are added as before.

*The solution of arsenic in dilute hydrochloric acid* requires about three grammes of acid carbonate of sodium, if 54 or 55 grammes of solution is the quantity employed. After boiling for a few minutes and cooling, the starch and iodine are added.

These arsenical solutions contain .9 per cent. of arsenic.

The *Hyposulphite of sodium* is dissolved in water, starch mucilage added, and the iodine solution slowly run in, the whole being frequently stirred, until a permanent blue color is produced.

In the previous reactions iodine has acted as an indirect oxidizing agent by uniting with the hydrogen and thus liberating the oxygen of water. In the present case it unites with an analogue of hydrogen, namely sodium, a new salt, tetrathionate of sodium, being simultaneously produced.

$$I_2 + 2(Na_2S_2O_3,5H_2O) = 2NaI + Na_2S_4O_6 + 10H_2O.$$

## QUESTIONS AND EXERCISES.

1003. Give equations illustrative of the reactions on which the use of a standard volumetric solution of iodine is based.

1004. From what point of view may iodine be regarded as an oxidizing agent?

1005. What reagent indicates the termination of the reaction between deoxidizing substances and moist iodine?

1006. How much sulphurous acid gas will cause the absorption of 2.54 parts of iodine in the volumetric reaction?—*Ans.* .64.

1007. What quantity of iodine will be required, under appropriate conditions, to oxidize 5 parts of arsenic?—*Ans.* 12.828.

1008. Find by calculation the amount of hyposulphite of sodium which will react with 13 parts of iodine in volumetric analysis.—*Ans.* 25.389.

---

### Standard Solution of Red Chromate of Potassium.

(Red Chromate of Potassium, $K_2CrO_4,CrO_3 = 295$.)

One molecule of red chromate of potassium in presence of an acid, under favorable circumstances, yields four atoms of oxygen to the hydrogen of the acid, leaving three available either for direct oxidation or for combination with the hydrogen of more acid, an equivalent proportion of acidulous radical being liberated for any required purpose.

When used as a volumetric agent, the red chromate always yields the whole of its oxygen to the hydrogen of the accompanying acid, a corresponding quantity of acidulous radicals being set free—four-sevenths of this radical immediately combining with the potassium and chromium of the red chromate, three-sevenths becoming available. Ferrous may thus be converted into ferric salts with sufficient rapidity and exactitude to admit of the estimation of an unknown quantity of iron by a known quantity of the red chromate. As one atom of the liberated acidulous radicals will convert two molecules of ferrous into one of ferric salts, one molecule of red chromate causes six of ferrous to become three of ferric, as shown in either of the following equations:—

$$K_2CrO_4,CrO_3 + 7H_2SO_4 + 6FeSO_4 = K_2SO_4,Cr_2SO_4 + 7H_2O + 3(Fe_2SO_4);$$

$$K_2CrO_4,CrO_3 + 14HCl + 6FeCl_2 = 2KCl + Cr_2Cl_6 + 7H_2O + 3Fe_2Cl_6.$$

These equations indicate that, in presence of excess of acid, 295 parts (the molecular weight) of red chromate of potassium will convert 1668 parts of crystallized ferrous sulphate. $6(FeSO_4,7H_2O = 278)$, or an equivalent quantity of ferrous chloride ($6FeCl_2 = 762$), ferrous carbonate ($6FeCO_3 = 696$), ferrous arseniate ($2Fe_2As_2O_8 = 892$), ferrous phosphate ($2Fe_2P_2O_8 = 716$), or iron itself ($3Fe_2 = 336$), into ferric salt. If these figures be taken to represent grammes, $\frac{1}{100}$ or less of the stated amounts will be found to be convenient quantities for experiment.

Dissolve 14.75 grammes of red chromate of potassium in one litre of distilled water. 100 c. c. of this solution contain $\frac{1}{100}$ of the molecular weight of the salt in grammes, and will cause the conversion of $\frac{1}{100}$ of 6 atomic weights of iron in grammes,

or an equivalent quantity of the lower salts of iron, from the ferrous to the ferric state. The solution is used in determining the strength of the following official ferrous preparations. It is known that the whole of the ferrous has been converted to ferric salt when a small drop of the liquid placed in contact with a drop of a very dilute solution of ferridcyanide of potassium, on a white plate, ceases to strike a blue color.

If the red chromate employed in making this standard solution is not known to be pure and dry, the strength of the solution may be checked by dissolving a small accurately weighed piece of piano-forte wire (0.4 or 0.5) in diluted sulphuric acid in a small flask, warmed, and then running in the solution of red chromate until conversion is effected. Every 1.68 part of iron used corresponds to 1.475 part of red chromate, and this weight of red chromate should be contained in 100 corresponding parts by volume of the standard solution. If the latter is stronger or weaker, water or salt respectively must be added; or else the strength or weakness must be noted and allowed for in every subsequent operation.

| | Grammes of substance. | | C. c. of vol. sol. |
|---|---|---|---|
| Iron, arseniate of . . . . . | 2.94 | = | 25 |
| " magnetic oxide of . . . . . | 2.00 | = | 8.3 |
| " phosphate of, B. P. and U. S. P. . | 2.00 | = | 25 |
| " saccharated carbonate of . . . . | 2.00 | = | 20.8 |

The several compounds are dissolved in excess of hydrochloric acid diluted with water, and the standard solution then dropped in. The ferrous liquid must not be exposed to the air for more than a few seconds after solution has been effected, or oxygen will be absorbed and a corresponding amount of ferric salt formed before the volumetric oxidizing agent is added; in most cases diluted sulphuric acid may be used as a solvent of the ferrous salt, ferrous sulphate absorbing oxygen from the air far less rapidly than ferrous chloride. It will be found that the proportion of carbonate of iron in the saccharated compound, as indicated by the above numbers, is, 37 in 100. Trade samples yield from 20 to 30, and sometimes 35 per cent., according to the care with which oxidation has been prevented. The theoretical percentage obtainable from the ingredients is 45.5, the quantity that would be present if the compounds were anhydrous and unoxidized, conditions never obtained in practice. Howie has suggested that as hydrochloric acid is known to so rapidly convert ordinary sugar into inverted sugar as to render it easily attacked by chromic acid, while phosphoric acid very slowly affects sugar, the latter acid instead of the

former should be employed in dissolving the saccharated carbonate of iron for volumetric analysis. Another mode of eliminating the influence of the sugar is to char it with oil of vitriol before analyzing. The 2 grammes of phosphate of iron should contain .895 of real ferrous phosphate or nearly 45 per cent. Trade samples often contain less, but sometimes more, than this. The magnetic oxide of iron (B. P.) is of official quality if it contain 9 per cent. of ferrous oxide. Theoretically it should contain 31 per cent. ($Fe_3O_4 = FeO + Fe_2O_3$ or $232 = 72 + 160$), but during desiccation in the air some oxidation occurs. The official figures for the arseniate of iron indicate about 38 per cent. of real arseniate, but commercial samples fall below even this low figure; in short, the compound is more nearly a ferric than a ferrous arseniate.

The use of these two volumetric solutions in quantitative analysis admits of great extension.

## QUESTIONS AND EXERCISES.

1009. Write equations explanatory of the oxidizing power of red chromate of potassium.

1010. One hundred cubic centimetres of an aqueous solution of red chromate of potassium contain $\frac{1}{197}$ of the molecular weight of the salt in grammes; with what weight of metallic iron, dissolved in hydrochloric acid, will this volume react?—*Ans.* 1.68 gramme.

1011. If 8.34 grammes of impure crystallized ferrous sulphate dissolved in acidulated water require 93 c. c. of the standard solution of chromate for complete conversion into ferric salt, what percentage of ferrous sulphate is present?—*Ans.* 93.

1012. How much red chromate of potassium is required for the conversion of 10 parts of ferrous sulphate into ferric salt?—*Ans.* 1.768.

1013. What quantity of pure ferrous carbonate is indicated by 1.475 part of red chromate as applied in volumetric analysis?—*Ans.* 3.48.

1014. State the amount of official saccharated carbonate of iron equivalent to .7375 part of red chromate in the volumetric reaction. —*Ans.* 4.7.

## ESTIMATION OF SUBSTANCES READILY DEOXIDIZED.

Any substance which quickly yields a definite amount of oxygen may be quantitatively tested by ascertaining how much of a deoxidizing agent of known power must be added to a given quantity before complete deoxidation is effected. The chief compounds which may be used as absorbers of oxygen (deoxidizers or reducing agents,

as they are commonly termed) are hyposulphite of sodium, sulphurous acid, ferrous sulphate,* oxalic acid, arsenious acid. The first-named is officially employed; it is only used in the estimation of free iodine, and, indirectly, of chlorine and chlorinated compounds. Iodine and chlorine are regarded as oxidizing agents, because their great affinity for hydrogen enables them to become powerful indirect oxidizers in presence of water.

### Standard Solution of Hyposulphite of Sodium.

(Crystallized Hyposulphite of Sodium, $Na_2S_2O_3,5H_2O = 248$.)

Dissolve about 30 grammes of hyposulphite of sodium in a litre or less of water. Fill a burette with this solution, and allow it to flow into a beaker containing exactly 100 c. c. of the volumetric solution of iodine until the brown color of the iodine is just discharged—or, starch being added, until the blue iodide of starch is decolorized. Note the number of c. c. of hyposulphite solution required, and to the bulk of the solution add water, so that 2.48 grammes of hyposulphite of sodium shall be contained in every 130 c. c.

When iodine and hyposulphite of sodium react, two atoms of iodine remove two of sodium from two molecules of the hyposulphite, tetrathionate of sodium being formed, as indicated in the following equation:—

$$I_2 + 2(Na_2S_2O_3,5H_2O) = 2NaI + Na_2S_4O_6 + 10H_2O.$$

As, therefore, 100 c. c. of the iodine solution contain $\frac{1}{100}$ of the atomic weight of iodine in grammes, 100 c. c. of the standard solution of hyposulphite of sodium will contain $\frac{1}{100}$ of the molecular weight of the salt in grammes, and will show the existence of $\frac{1}{100}$ of the atomic weight of iodine in grammes in any quantity of a liquid normally containing free iodine, or iodine liberated by an equivalent quantity of free chlorine.

This solution is employed for quantitatively testing the following substances:—

| | Grammes of substance. | | C. c. of vol. sol. |
|---|---|---|---|
| Chlorine, solution of . . . . . | 29.26 | = | 50 |
| Iodine, B. P. and U. S. P. . . . . | 1.27 | = | 100 |
| Lime, chlorinated . . . . . | 1.17 | = | 100 |
| " solution of chlorinated . . . | 6.00 | = | 50 |
| Soda, solution of chlorinated . . . | 7.00 | = | 50 |

* "Five grains of Permanganate of Potassium dissolved in water require for decoloration a solution of forty-four grains of granulated sulphate of iron acidulated with two fluidrachms of diluted sulphuric acid."—B. P.

*Note.*—Owing to the volatility of chlorine and iodine, and the readiness with which they attack the metals of which balances are made, it is not desirable to experiment on stated weights of substances containing these elements. A small stoppered bottle or tube containing the material may be counterpoised, and a convenient quantity removed for analysis, the precise amount taken being ascertained by again weighing the bottle.

*The iodine* may be dissolved in water containing about a gramme and a half of iodide of potassium, a salt giving no reaction with hyposulphite of sodium.

$$\underbrace{I_2}_{254} \text{ eq. to } \underbrace{2(Na_2S_2O_3,5H_2O)}_{496}$$

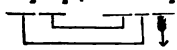
It is assumed in this operation that the iodine has been shown by qualitative analysis to be free from chlorine and bromine. These elements resemble iodine in reacting with hyposulphite of sodium, hence would reckon as iodine in a volumetric assay.

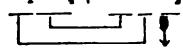
*The solution of chlorine* is added to water containing excess of iodide of potassium (about a gramme and a third); a quantity of iodine, equivalent to the amount of chlorine present, is thus liberated. The hyposulphite solution is then dropped in.

$$\underbrace{Cl_2}_{71} \text{ eq. to } \underbrace{2(Na_2S_2O_3,5H_2O}_{496}$$

*The chlorinated lime* ("chloride of lime" or "bleaching-powder") is mixed with about a fifth of a litre of water containing excess of iodide of potassium (3.5 grms.) and acidulated with hydrochloric acid. Diluted sulphuric acid would act in a similar manner:—

$H_2Cl_2 + CaOCl_2$ $\qquad$ $H_2SO_4 + CaOCl_2$





The chlorine first set free liberates an equivalent amount of iodine from the iodide of potassium, $Cl_2 + 2KI = 2KCl + I_2$. The hyposulphite and iodine reacting show the value or the indirect oxidizing power of the chlorinated lime; it should correspond to 30 per cent. of chlorine. For example, as 248 (1 molecule) of hyposulphite indicate the presence of 35.5 (1 atom) of chlorine, 2.48 of hyposulphite (the quantity in 100 c. c.) indicate the presence of .355 of chlorine, if 100 c. c.

have been used, therefore, .355 of chlorine is obtainable from the quantity of bleaching-powder employed (1.17). And if 1.17 of bleaching-powder yield .355 of chlorine, 100 will yield about 30 (30$\frac{1}{2}$ nearly).

*The solution of chlorinated lime* is mixed with about a fifth of a litre of water containing a couple of grammes of iodide of potassium, and ten or twelve c. c. of hydrochloric acid. The hyposulphite solution is then added from a burette until the color of the liberated iodine is just discharged. The solution of chlorinated soda is similarly treated.

*Note.*—1. In these experiments the blue color formed on the addition of mucilage of starch to the liquids will be found to be a more delicate indicator of the termination of reactions than the brown tint of the iodine.

*Note.*—2. Standard solutions used in volumetric analysis are often described as normal, decinormal, and centinormal. A normal solution (N) contains in every litre the molecular weight of the salt, *taken in grammes;* a decinormal solution is one-tenth ($\frac{N}{10}$), and a centinormal ($\frac{N}{100}$) one-hundredth the strength of such a normal solution.

## QUESTIONS AND EXERCISES.

1015. For what purposes is the official volumetric solution of hyposulphite of sodium used?

1016. On what reaction is based the quantitative employment of hyposulphite of sodium?

1017. How much hyposulphite of sodium is required to show the presence of 10 parts of iodine?—*Ans.* 19.527.

1018. To what amount of chlorine is 4.96 parts of hyposulphite of sodium equivalent in volumetric analysis?—*Ans.* 0.71.

1019. Describe the operation included in the estimation of the strength of bleaching powder.

1020. By what reagent is the complete absorption of free iodine by hyposulphite of sodium indicated?

## MISCELLANEOUS PROBLEMS.

1021. How much bicarbonate of potassium is contained in an eight-ounce bottle of medicine, seven fluidrachms of which are saturated by two and a half grains of crystallized oxalic acid?—*Ans.* 36.3 grains.

1022. A sample of soda-ash is said to contain 78 per cent. of pure anhydrous carbonate of sodium; if the statement is true, how much of the official volumetric solution of oxalic acid will saturate 5 grammes of the specimen?—*Ans.* 73.6.

1023. 2.69 grammes of common brown sulphuric acid are saturated by 43.5 cubic centimetres of the official volumetric solution of soda; how much acid of 96.8 per cent. is present?—*Ans.* The 2.69 contained 2.2.

1024. Four grammes of a litre and a half of concentrated hydrocyanic acid are neutralized by 89 cubic centimetres of volumetric solution of nitrate of silver of official strength: to what volume must the bulk of the acid be diluted for the production of acid of pharmacopœial strength?—*Ans.* 9 litres.

1025. 3.18 grammes of a powder containing arsenic require for complete reaction 84 cubic centimetres of a volumetric solution of iodine, which is 1.43 weaker than the standard solution of the British Pharmacopœia; what percentage of pure arsenic is contained in the powder?—*Ans.* 12.86.

1026. How much pure metal is present in a sample of iron 1.68 gramme of which, dissolved in dilute sulphuric acid, is exactly attacked by 95.7 cubic centimetres of semi-decinormal volumetric solution of red chromate of potassium which is 6 per cent. too strong?

---

# GRAVIMETRIC QUANTITATIVE ANALYSIS.

## ESTIMATION OF METALS.

### POTASSIUM.

*Outline of the Process.*—This element is usually estimated in the form of double chloride of potassium and platinum. Qualitative analysis having proved the presence of potassium and other radicals in a substance, a small quantity of the material is accurately weighed, dissolved, and the other elements removed by appropriate reagents; the precipitates are well washed, in order that no trace of the potassium salt shall be lost, the resulting liquid concentrated over a water-bath (to avoid loss that would occur mechanically during ebullition), hydrochloric acid added if necessary, solution of perchloride of platinum poured in, and evaporation continued to dryness; excess of the perchloride is then dissolved out by adding, to the dried residue, spirit of wine containing half its bulk of ether (a liquid in which the double chloride is insoluble), the mixture carefully poured on to a tared and dried filter, washed with the spirit till every trace of free perchloride of platinum is removed, the whole dried and weighed; from the resulting amount the proportion of potassium, or equivalent quantity of a salt of potassium, is ascertained by calculation.

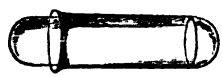
*Note.*—From this short description it will be seen, first, that the chemistry of quantitative is the same as that of qualitative analysis; the second, that the principle of gravimetric is the same as that of volumetric quantitative analysis—the combining proportions being known, unknown quantities of elements may be ascertained by calculation from known quantities of their compounds.

*Apparatus.*—In addition to a delicate balance and weights and the common utensils, a few special instruments are used in quantitative manipulation; some of these may be prepared before proceeding with the estimation of potassium.

*Filtering-paper* may be of the kind known as "Swedish," the texture of which is of the requisite degree of closeness, and its ash small in amount. A large number of circular pieces of one size, six to eight centimetres in diameter, should be cut ready for use. In delicate experiments, where a precipitate on a filter has to be ignited and the paper consequently burnt, the weight of the ash of the filter must be deducted from the weight of the residue. The ash is estimated after burning ten or twenty of the cut filters. These are folded into a small compass, a portion of a piece of platinum wire twisted a few times round the packet, so as to form a cage, the whole held by the free end of the wire over a weighed porcelain crucible placed in the centre of a sheet of glazed paper, the bundle ignited by a spirit-lamp or smokeless gas-flame, the flame allowed to impinge against the charred mass till it falls into the crucible below, any stray fragments on the sheet carefully shaken into the crucible, the latter placed over a flame till carbon has all burnt off and nothing but ash remains, the whole cooled, weighed, and the weight of the crucible deducted; the weight of the residue divided by the number of pieces used give the average amount of ash in each filter.

*A pair of weighing-tubes* (Fig. 74), for holding dried filters during operations at the balance, may be made from two test-

Fig. 74.



A pair of weighing-tubes.

Fig. 75.

Clamped watch-glass for weighing.

tubes, one fitting closely within the other. About five centimetres of the closed end of the outer and seven of the inner are cut off, by leading a crack round the tube with a pencil of incandescent charcoal, and the sharp edges fused in

the blowpipe-flame. A filter, after drying, is quickly folded and placed in the narrower tube, the mouth of which is then closed by the wider tube. This prevents reabsorption of moisture from the air.

*A pair of Watch-glasses*, having accurately ground edges and clamped, as shown in Fig. 75, also forms a convenient arrangement for weighing filters, etc.

*The Washing-bottle*, holding the *spirit of wine and ether*, is a common flask, through the cork of which a short straight tube passes. The outer end of the tube should be sufficiently narrowed to enable it to deliver a very fine stream of the liquid. The flask being inverted, the warmth of the hand expands the air and vapor to a sufficient extent to force out the liquid.

Fig. 76.

The washing-bottle.

*The ordinary Washing-bottle* for quantitative operations should be formed of a flask in which water may be boiled, fitted up as usual (*vide* p. 107).

*A Water-oven* is the best form of drying-apparatus. It is a small square copper vessel, jacketed on five sides and having a door on the sixth; water is poured into the space between the inner and outer casing, and the whole placed over a gas-lamp or source of heat, moist air and steam escaping by appropriate apertures. Desiccation at higher temperatures than the boiling-point of water may be practised by using oil or paraffin instead of water, inserting a thermometer in the fat. The apparatus may be purchased of any maker of chemical instruments.

*Pure distilled water* must be used in all quantitative determinations.

*Note.*—In practising the operations of quantitative analysis, experiments should at first be conducted on definite salts of known composition, for the accuracy of results may then be tested by calculation.

*Estimation of Potassium in the form of double coloride of potassium and platinum.*—Select two or three crystals of pure nitrate of potassium, powder them in a clean mortar, dry the powder by gently heating in a porcelain crucible over a flame for a few seconds, place about a couple of decigrammes (0.2 grm.) of the powder in a counterpoised watch-glass, accurately weigh the selected quantity, transfer to a small dish, letting water from a wash-bottle flow over the watch-glass and run into the dish, warm the dish till the nitrate is dissolved, acidulate with hydrochloric acid, add excess of aqueous solution of perchloride of platinum (a quantity containing about 0.4 of

solid salt), evaporate to dryness over a water-bath. While evaporation is going on, place a filter and the weighing-tubes in the water-oven, exposing them to a temperature of 100° C. for about half an hour; fold the filter and insert it in the tubes, place them on a plate under a glass shade, and when cold, accurately note their weight. Arrange the weighed filter in a funnel over a beaker. Transfer the dried and cooled platinum salt from the dish to the filter by moistening the residue with the mixture of alcohol and ether, and, when the salt is loosened, pouring the contents of the dish into the paper cone. Any salt still adhering may be freed by the finger, which, together with the dish, should be washed in the stream of spirit, the rinsings at once flowing into the filter. The filtrate should have a yellowish-brown color, due to the excess of perchloride of platinum. If it is colorless, an insufficient amount of perchloride has been added, and the whole operation must be repeated. The washed precipitate and filter are finally dried in the water-oven, folded and placed in the weighing-tubes, the drying continued until the whole, after repeated weighing when cold, ceases to alter; the final weight is noted.

*Note.*—If filters are not freed from all trace of acid by thorough washing, the paper will be brittle when dry, falling to pieces on being folded.

*Analytical memoranda* in the note book may have the following form:—

| | |
|---|---|
| Watch-glass and substance . . . . . | |
| Watch-glass . . . . . | |
| Substance . . . . | |
| Weighing-tubes, filter, and Pt salt . . . | |
| Weighing-tubes and filter . . . . . | |
| $PtCl_4,2KCl$ . . . | |

The calculations are simple:—

$$\text{As } \left\{\begin{matrix}PtCl_4,2KCl\\ =189\end{matrix}\right\} \text{ are equivalent to } \left\{\begin{matrix}2KNO_3\\ =202\end{matrix}\right\}$$

$$\text{so } \left\{\begin{matrix}\text{the weight of}\\ \text{double chloride}\\ \text{obtained}\end{matrix}\right\} \text{ is equivalent to } x.$$

$x$ will be the amount of *pure* nitrate of potassium in the quantity of substance operated on. $x$ should, in the present instance, be identical with the weight of substance taken, because, for educational purposes, pure nitre is under examination. Only after analyses of

pure substances have yielded the operator results identical with those by calculation, can analyses of substances of unknown degree of purity be undertaken with confidence. A table of atomic weights, from which to find molecular weights, is given in the Appendix.

*Platinum residues* should be preserved, and the metal recovered from them from time to time (*vide* p. 248).

Hot alcohol sometimes reduces perchloride of platinum, the metal being thrown out of solution in a finely divided form, known as *platinum black;* only aqueous solutions, therefore, of the salt should be used where heat is employed. Hence, also, in washing out excess of perchloride of platinum from the double chloride of platinum and potassium by spirit, the application of heat should be avoided.

*Effervescing Potash-water* (*Liquor Potassæ Effervescens*, B. P.) is most easily estimated volumetrically (p. 550). Any adulteration by an equivalent amount of bicarbonate of sodium would, however, by that process be undetected; hence the Pharmacopœia directs that "five fluidounces, evaporated to one-fifth, and 12 grains of tartaric acid added, yield a crystalline precipitate, which, when dried, weighs not less than 12 grains." Five fluidounces of this preparation should contain 7.5 grains of bicarbonate, convertible into 14.1 grains of acid tartrate of potassium by 11.25 grains of tartaric acid. The method is somewhat rough, but quite efficient for "potash-water" containing nothing but bicarbonates of alkali-metals.

*Proportional weights of equivalent quantities of potassium and its salts.*

| | | |
|---|---|---|
| Metal | $K_2$ | 78 |
| Oxide ("potash") | $K_2O$ | 94 |
| Hydrate ("caustic postash") | $2KHO$ | 112 |
| Carbonate (anhydrous) | $K_2CO_3$ | 138 |
| Carbonate (crystalline) | $K_2CO_3 + 16$ % aq. | 164.285 |
| Bicarbonate | $2KHCO_3$ | 200 |
| Nitrate | $2KNO_3$ | 202 |
| Platinum salt | $PtCl_4,2KCl$ | 489 |

## SODIUM.

Sodium is usually estimated as sulphate. Accurately weigh a porcelain crucible and lid, place within about .3 of pure rock-salt, and again weigh, making a memorandum of the weights in a note-book. Add rather more strong sulphuric acid than may be considered sufficient to convert the chloride into acid sulphate of sodium. Heat the crucible gradually, the flame being first directed against the side of the crucible to avoid violent ebullition, until fumes of acid cease to be evolved,

towards the end of the operation dropping in one or two fragments of carbonate of ammonium to facilitate complete expulsion of all excess of acid. When cold weigh the crucible and contents. The weight of the crucible having been deducted, the amount of sulphate obtained should be the exact equivalent of the quantity of chloride of sodium employed.

$$\underbrace{2NaCl}_{117} + H_2SO_4 = \underbrace{Na_2SO_4}_{142} + 2HCl.$$

*Proportional weights of equivalent quantities of sodium and its salts.*

| | | |
|---|---|---|
| Metal | $Na_2$ | 46 |
| Oxide ("soda") | $Na_2O$ | 62 |
| Hydrate ("caustic soda") | $2NaHO$ | 80 |
| Carbonate (anhydrous) | $Na_2CO_3$ | 106 |
| Carbonate (crystals) | $Na_2CO_3,10H_2O$ | 286 |
| Bicarbonate | $2NaHCO_3$ | 168 |
| Chloride | $2NaCl$ | 117 |
| Sulphate (anhydrous) | $Na_2SO_4$ | 142 |
| Sulphate (crystals) | $Na_2SO_4,10H_2O$ | 322 |

## AMMONIUM.

Salts of ammonium are, for purposes of quantitative analysis, generally converted into the double chloride of ammonium and platinum ($PtCl_4 2NH_4Cl$), the details of manipulation being the same as those observed in the case of potassium. About 0.15 grm. of pure, white, dry chloride of ammonium may be taken for experiment.

COMPOSITION OF THE PLATINUM SALT.

| | | In 1 molecule. | In 100 parts. |
|---|---|---|---|
| Pt | 198 | 198 | 44.30 |
| $Cl_6$ | $35.5 \times 6$ | 213 | 47.64 |
| $N_2$ | $14.0 \times 2$ | 28 | 6.27 |
| $H_8$ | $1.0 \times 8$ | 8 | 1.79 |
| | | 447 | 100.00 |
| or, $PtCl_4$ | 340 | 340 | 76.06 |
| $2NH_4Cl$ | $53.5 \times 2$ | 107 | 23.94 |
| | | 447 | 100.00 |

The proportion of nitrogen, ammonium, or chloride of ammonium in the double chloride may also be ascertained from the weight of platinum left on igniting the double chloride; for this purpose heat must be applied slowly, or platinum will be mechanically carried off with the gaseous products of decomposition.

*Proportional weights of equivalent quantities of ammoniacal compounds.*

| | | |
|---|---|---|
| Ammonia (gas) . . . . . | $2NH_3$ . . . . . | 34 |
| Ammonium . . . . . | $(NH_4)_2$? . . . . | 36 |
| Chloride of ammonium . . . | $2NH_4Cl$ . . . . | 107 |
| Platinum salt . . . . . | $PtCl_4,2NH_4Cl$ . . | 447 |
| "Carbonate of ammonium" . . | $(N_4H_{16}C_3O_8) \div 2$ . | 118 |
| Sulphate of ammonium . . . | $(NH_4)_2SO_4$ . . . | 132 |

## BARIUM.

Barium is estimated in the form of anhydrous sulphate of barium ($BaSO_4$).

*Process.*—Dissolve 0.3 or 0.4 of pure crystallized and dried chloride or nitrate of barium in about half a litre of water in a beaker, heating to incipient ebullition, and slightly acidulating with hydrochloric or nitric acid. Add diluted sulphuric acid (prepared some days previously, so that sulphate of lead may have deposited) so long as a precipitate forms, keep the mixture hot for some time, set aside for half an hour, pass the supernatant liquid through a filter, gently boil the residue two or three times with more water; finally collect the precipitate on the filter, removing adherent particles from the beaker by the finger, and cleansing by a stream of hot water from the wash-bottle. The precipitate must be washed with hot water until the filtrate ceases to turn litmus-paper red, or give any cloudiness when tested with chloride of barium. The filter and sulphate of barium, having thoroughly drained, is dried in a warm place, commonly by supporting the funnel in an inverted bottomless beaker over a sand-bath or hot plate.

The sulphate of barium is now removed from the filter, heated to drive off every trace of moisture, and weighed. This is accomplished by placing a weighed porcelain crucible (and cover) on a sheet of glazed paper, holding the filter over it, and carefully transferring the precipitate; the sides of the filter are then gently rubbed together and detached powder dropped into the crucible, the paper folded, encased in two or three coils of one end of a platinum wire and burnt over the

crucible, ash and any particles in the sheet of paper dropped into the sulphate of barium, the open crucible exposed over a flame till its contents are quite white, covered, cooled, and weighed.

| | Formula. | Molecular weights. |
|---|---|---|
| Chloride of barium | $BaCl_2 2H_2O$ | 244 |
| Nitrate of barium | $Ba2NO_3$ | 261 |
| Sulphate of barium | $BaSO_4$ | 233 |

*Composition of Sulphate of Barium.*

| | | In one molecule. | In 100 parts. |
|---|---|---|---|
| Ba | 137 | 137 | 58.80 |
| S | 32 | 32 | 13.73 |
| $O_4$ | 16×4 | 64 | 27.47 |
| | | 233 | 100.00 |

In these educational experiments it is unnecessary to take filter-ash into account. Inevitable mistakes of manipulation commonly cause far greater errors.

## CALCIUM.

Calcium is usually thrown out of solution in the form of oxalate, the precipitate ignited, and the resulting carbonate weighed.

*Process.*—Dissolve 0.3 or 0.4 of dried colorless crystals of calc-spar in about a third of a litre of water acidulated with hydrochloric acid, heat the solution to near the boiling-point, add excess of solution of oxalate of ammonium, then ammonia until, after stirring, the liquid smells strongly ammoniacal; set aside in a warm place for twelve hours. Carefully pour off the supernatant liquid, passing it through a filter; add hot water to the precipitate, set aside for half an hour, again decant, and after once more washing, transfer the precipitate to the filter, allowing all contained fluid to pass through before a fresh portion is added. Wash the precipitate with hot water, avoiding a rapid stream, or the precipitate may be driven through the pores of the paper. Dry, transfer to a weighed crucible, and incinerate, as described for sulphate of barium, and slowly heat the precipitate till the bottom of the crucible is just visibly red when seen in the dark. As soon as the residue is white, or only faintly gray, remove the lamp, cool, and weigh.

The resulting carbonate of calcium should have the same weight as the calc-spar from which it was obtained. If loss has occurred, carbonate acid gas has probably escaped. In that

case moisten the residue with water, and after a few minutes test the liquid with red litmus or turmeric paper; if an alkaline reaction is noticed, it is due to the presence of caustic lime. Add a small lump of carbonate of ammonium, evaporate to dryness over a water-bath, and again ignite, this time being careful not to go beyond the prescribed temperature. The treatment may, if necessary, be repeated.

*Proportional weights of equivalent quantities of calcium salts.*

| | | |
|---|---|---|
| Oxide (quicklime) | CaO | 56 |
| Hydrate (slaked lime) | Ca2HO | 74 |
| Carbonate | $CaCO_3$ | 100 |
| Sulphate (anhydrous) | $CaSO_4$ | 136 |
| Sulphate (crystalline or precipit'd) | $CaSO_4,2H_2O$ | 172 |
| Chloride | $CaCl_2$ | 111 |
| Phosphate (of bone) | $(Ca_32PO_4)310 \div 3$ | 103.3 |
| Superphosphate | $CaH_42PO_4$ | 234 |

## MAGNESIUM.

*Process* 1.—*The light or heavy carbonate of magnesium* of pharmacy may be estimated by heating a weighed quantity to redness in a porcelain crucible. If it has the composition indicated by the formula given in the British Pharmacopœia ($3MgCO_3,Mg2HO,4H_2O$), it will yield 42 per cent. of magnesia (MgO). According to that work, the purity of even sulphate of magnesium ($MgSO_4,7H_2O$) may be determined by boiling a weighed quantity with excess of carbonate of sodium, collecting the precipitate, washing, drying, igniting, and weighing the resulting magnesia (MgO). The crystalline sulphate should afford 16.26 per cent. of oxide. The official solution of carbonate of magnesium in carbonic acid water (*Liquor Magnesiæ Carbonatis*, B. P.) should yield five grains of pure oxide of magnesium per fluidounce.

*Process* 2. — *The general form in which magnesium is precipitated* is as phosphate of ammonium and magnesium ($MgNH_4PO_4,6H_2O$); this, by heat, is converted into pyrophosphate of magnesium ($Mg_2P_2O_7$). Accurately weigh a small quantity (0.4 to 0.5) of pure dry crystals of sulphate of magnesium, dissolve in two or three hundred cubic centimetres of cold water in a beaker, add chloride of ammonium, ammonia, and phosphate of sodium or ammonium, agitate with a glass rod (without touching the sides of the vessel, or crystals will firmly adhere to the rubbed portions) and set aside for twelve hours.

Collect on a filter, wash the precipitate with water containing a tenth of its volume of the strongest solution of ammonia, until the filtrate ceases to give a precipitate with an acidulated solution of nitrate of silver. Dry, transfer to a crucible, burn the filter in the usual way, heat slowly to redness, cool, and weigh.

*Proportional weights of equivalent quantities of magnesium salts.*

| | | |
|---|---|---|
| Pyrophosphate . . | $Mg_2P_2O_7$ . . . . . | 222 |
| Sulphate . . . . | $2(MgSO_4,7H_2O)$ . . . . . | 492 |
| Oxide . . . . . | $2(MgO)$ . . . . . | 80 |
| Official carbonate . | $(3MgCO_3,Mg2HO,4H_2O)\div 2$ . | 191 |

## ZINC.

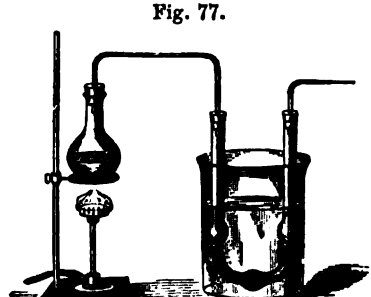
Zinc is usually estimated as oxide (ZnO), occasionally as sulphide (ZnS).

*Process.*—Dissolve a weighed quantity (0.5 to 0.6) of sulphate of zinc in about half a litre of water in a beaker, heat to near the boiling-point, add carbonate of sodium in slight excess, boil, set aside for a short time; pass the supernatant liquid through a filter, gently boil the precipitate with more water, again decant; repeat these operations two or three times; collect the precipitate on the filter, wash, dry, transfer to a crucible, incinerate, ignite, cool, and weigh. 287 (= molec. weight) of sulphate should yield 81 (= molec. weight) of oxide.

## MANGANESE.

To ascertain its value for evolving chlorine from hydrochloric acid a weighed quantity of finely powdered black oxide of

Fig. 77.



manganese is heated in a small flask with pure hydrochloric acid, and the resulting chlorine conveyed into a U-tube containing solution of iodide of potassium. The amount of iodine thus freed is estimated by the volumetric solution of hyposulphite of sodium. 127 of iodine indicate 35.5 of chlorine. Manganese may also be estimated by the reaction and apparatus described under "Oxalates," page 592.

## ALUMINIUM.

Aluminium is always precipitated as hydrate ($Al_2 6HO$) and weighed as oxide ($Al_2O_3$).

*Process.*—Dissolve about two grammes of pure dry ammonium-alum in half a litre of water, heat the solution, add chloride of ammonium and a slight excess of ammonia, boil gently till the odor of ammonia has nearly disappeared, set aside for the hydrate to deposit, pass the supernatant liquid through a filter, wash the precipitate three or four times by decantation, transfer to the filter, finish the washing, dry, burn the filter, ignite in a covered crucible, and weigh.

| | |
|---|---|
| $Al_2 3SO_4, (NH_4)_2SO_4, 24H_2O$ . . . . . | 907 |
| $Al_2O_3$ . . . . . | 103 |
| Per cent. of $Al_2O_3$ yielded by ammonium-alum . . | 11.356 |

---

## QUESTIONS AND EXERCISES.

1027. Give details of the manipulations observed in gravimetrically estimating salts of potassium or ammonium.

1028. What quantity of chloride of sodium is contained in a sample of rock-salt 0.351 gramme of which yields 0.426 of sulphate of sodium? —*Ans.* 100 per cent. (It is absolutely pure.)

1029. To what amount of the official alum is 0.894 of a gramme of the double chloride of platinum and ammonium equivalent?—*Ans.* 1.814 gramme.

1030. Find the weight of sulphate of barium obtainable from 0.522 of nitrate.—*Ans.* 0.466.

1031. Describe the usual method by which salts of calcium are estimated.

1032. By what quantitative processes may the official salts of magnesium be analyzed?

1033. Calculate the proportion of pure sulphate of zinc in a sample of crystals 0.574 of which yield 0.161 of oxide.—*Ans.* 99.3 per cent.

1034. Ascertain the weight of alumina ($Al_2O_3$) which should be obtained from 1.814 gramme of ammonium-alum.

## IRON.

Iron and its salt are gravimetrically estimated in the form of ferric oxide ($Fe_2O_3$).

*Compounds containing organic acidulous radicals* are simply incinerated, and the resulting oxide weighed. Thus 1 gramme of the official citrate of iron and ammonium (*Ferri et Ammoniæ Citras*, B. P.) incinerated, with exposure to air, leaves not less than .27 of ferric oxide. A small quantity of the salt is weighed in a tared covered porcelain crucible, flame cautiously applied until vapors cease to be evolved, the lid then removed, the crucible slightly inclined and exposed to a red heat until all carbonaceous matter has disappeared. The residual ferric oxide is then weighed. The tartrate of potassium and iron (*Ferrum Tartaratum*, B. P.) is treated in the same manner, except that the ash must be washed and again heated before weighing, in order to remove carbonate of potassium produced during incineration; 5 grammes should yield 1.5 gramme of ferric oxide.

*From other compounds* of iron, soluble in water or acid, the metal is precipitated in the form of hydrate ($Fe_2 6HO$) by solution of ammonia, and converted into oxide ($Fe_2O_3$) by ignition. Dissolve a piece (about 0.2) of the purest iron obtainable (piano wire), accurately weighed, in water acidulated with hydrochloric acid; add a few drops of nitric acid and gently boil; pour in excess of ammonia, stir, set aside till the ferric hydrate has deposited, pass the supernatant liquid through a filter, treat the precipitate three or four times with boiling water; transfer to the filter, wash till the filtrate yields no trace of chlorine (for chloride of ammonium will decompose ignited ferric oxide, with volatilization of ferric chloride), dry and ignite as usual and weigh. Iron in the official solutions (*Liquor Ferri Perchloridi Fortior*, *Liquor Ferri Pernitratis*, and *Liquor Ferri Persulphatis*) may be estimated by this general process.

*The proportion of metallic iron in a mixture* of iron and oxides of iron may be determined by digestion in a strong solution of iodine in iodide of potassium, which attacks the metal only. The reduced iron of pharmacy (*Ferrum Redactum*) is in good condition so long as it contains, as shown by this method, half its weight of free metal.

*Proportional weights of equivalent quantities of iron and its salts.*

| | | |
|---|---|---|
| Metal | $Fe_2$ | 112 |
| Ferric oxide | $Fe_2O_3$ | 160 |
| Ferric hydrate | $Fe_26HO$ | 214 |
| Ferric chloride | $Fe_2Cl_6$ | 325 |
| Ferric sulphate | $Fe_23SO_4$ | 400 |
| Ferrous sulphate | $2(FeSO_4,7H_2O)$ | 556 |

## ARSENICUM.

Arsenic ($As_2O_3$) is usually estimated volumetrically (*vide* p. 560). With certain precautions arsenicum may also be precipitated and weighed as sulphide ($As_2S_3$).

*Process.*—The pure, white, massive arsenic (about 0.2) is dissolved in a flask in a small quantity of water containing bicarbonate of sodium or potassium, the liquid being heated. A slight excess of hydrochloric acid is then added, and sulphuretted hydrogen gas passed through the solution so long as a precipitate falls, the mouth of the flask being stopped by a plug of cotton-wool (to prevent undue access of air and consequent decomposition of the gas, resulting in precipitation of sulphur). The mixture is warmed in the flask and carbonic acid gas passed through it until the odor of sulphuretted hydrogen has nearly disappeared; the precipitate collected on a tared filter, washed as quickly as possible with hot water containing a little sulphuretted hydrogen, dried in a water-oven, and weighed. 198 parts of arsenic should yield 246 of sulphide of arsenicum.

## ANTIMONY.

The metal is precipitated in the form of sulphide ($Sb_2S_3$), with the precautions observed in estimating arsenicum—a small quantity of tartaric acid, as well as hydrochloric, being added, to prevent the precipitation of an oxysalt. If the sulphuretted hydrogen be passed through a hot solution, the particles of precipitate aggregate better, and the latter may be more quickly filtered out and washed. The experiment may be performed on about half a gramme of pure tartar-emetic: the salt should yield nearly half its weight (49.56 per cent.) of sulphide. According to Fresenius, the sulphide dried at 100° C. still contains 2 per cent. of water, and must be heated, in a current of carbonic acid gas, until it turns from an orange to a black color, before all moisture is expelled. In the British

Pharmacopœia the purity of tartar-emetic (*Antimonium Tartaratum*), and the strength of solution of chloride of antimony (*Liquor Antimonii Chloridi*), are determined by the above process.

## COPPER.

Copper is precipitated from its solutions and weighed (1) as metal ($Cu_2$), or (2) as oxide (CuO).

*Process* 1.—Dissolve about half a gramme of dry crystallized sulphate of copper in a small quantity of water, in a tared porcelain crucible or beaker, acidulate with hydrochloric acid, introduce a fragment or two of pure zinc, cover the vessel with a watch-glass, and set aside till evolution of hydrogen has ceased and the still acid liquid is colorless. The copper is then washed with hot water by decantation until no trace of acid remains, the precipitate drained, rinsed with strong spirit of wine, dried in the water-oven, and weighed.

*Process* 2.—About three-fourths of a gramme of sulphate of copper is accurately weighed, dissolved in half a litre of water, the liquid boiled; dilute solution of potash or soda is then added till no more precipitate falls, ebullition continued for a short time, and the beaker set aside; the supernatant liquid is decanted, the precipitate boiled with water, twice or thrice collected on a filter, washed, dried, transferred to a crucible, the filter incinerated, and its ash moistened with a drop of nitric acid; the whole is finally heated strongly, cooled, and weighed.

249.5 parts of sulphate of copper yield 79.5 of oxide, or 63.5 of metal.

## BISMUTH.

Dissolve 0.3 or 0.4 of pure oxycarbonate of bismuth ($2Bi_2O_2CO_3.H_2O$) (*Bismuthi Carbonas*, B. P.) in a small quantity of hydrochloric acid, dilute with water slightly acidulated by hydrochloric acid, pass excess of sulphuretted hydrogen through the liquid, collect the precipitate on a tared filter, wash, dry at 100° C., and weigh. The sulphide must not be exposed too long in the water-oven, or it will increase in weight owing to absorption of oxygen; hence it should be tested in the balance every half-hour during desiccation. 517 of oxycarbonate should yield 512 of sulphide ($Bi_2S_3$). The strength of the official solution of citrate of bismuth and ammonium (*Liquor Bismuthi et Ammoniæ Citratis*, B. P.) is determined by this process. "Three fluidrachms of the solution,

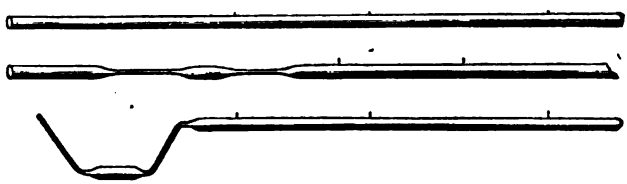
mixed with an ounce of distilled water, and treated with sulphuretted hydrogen in excess, yield a black precipitate, which collected, washed, and dried, weighs 9.92 grains. One fluidrachm yields three grains of oxide of bismuth." The atomic weight of bismuth is 208.

## MERCURY.

This element may be (1) isolated and estimated in the form of metal, or precipitated and weighed as (2) mercurous chloride, or (3) mercuric sulphide.

*Process* 1.—The process by which the metal itself is separated is one of distillation, into a bulb surrounded by water.

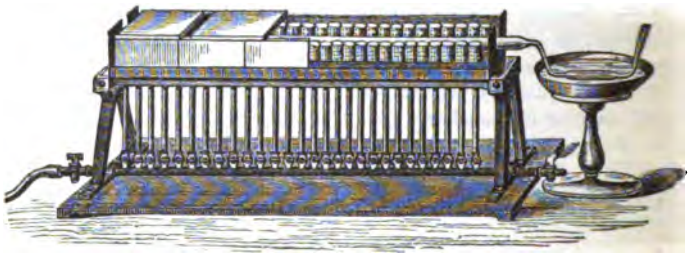
Figs. 78, 79, 80.



About half a metre of the difficultly fusible German glass known as *combustion-tubing* is sealed at one end after the manner of a test-tube (Fig. 78); a mixture of bicarbonate of sodium and dry chalk is then dropped into the tube to the height of two or three centimetres, and, next, several small fragments of quicklime so as to occupy another centimetre; a mixture of about a gramme of pure calomel or corrosive sublimate with enough powdered quicklime to occupy 10 or 12 centimetres of the tube is added, then the lime-rinsings of the mixing-mortar, a layer of a few centimetres of powdered quicklime, and finally a plug of *asbestos* (a fibrous mineral unaffected by heat). The whole powder should occupy two-thirds of the length of the tube. The part of the tube just above the asbestos is now softened in the blowpipe-flame and drawn out about a decimetre to the diameter of a narrow quill; it is again drawn out to the same extent at a point about two or three centimetres nearer the mouth, and any excess of tubing cut off. The bulb thus formed may be enlarged by softening and blowing. The tube is next softened at a point close to but anterior to the asbestos, and bent to form an obtuse angle; the tube is then softened close to the bulb and slightly bent so that the bulb may be parallel with the large tube; then

softened on the other side of the bulb, and the terminal tube bent to an obtuse angle, so that, the tube being held in a horizontal position, the bulb may be sunk in water, and the terminal tube point upwards (Fig. 80). The long tube is now laid in the gas-furnace found in most laboratories (Fig. 81), a basin so placed that the bulb of the apparatus may be cooled by

Fig. 81.



Distillation of Mercury for Quantitative Purposes.

being surrounded by water, the part of the tube occupied by asbestos heated to redness, and the flame slowly lengthened until the whole tube is red-hot. Under the circumstances just described the mercurial compound volatilizes, is decomposed by the lime, and its acidulous radical fixed, the mercury carried to and condensed in the bulb, the carbonic acid gas evolved from the bicarbonate of sodium and chalk washing out the last portions of mercury-vapor from the tube. When the distillation is considered to be complete, the dish of water is removed, the bulb dried, and then detached by help of a file at a point beyond any sublimate of mercury. The bulb is lastly weighed, the mercury shaken, or dissolved out, and the tube again dried and weighed.

*Process* 2.—The process by which mercury is separated in the form of calomel consists in adding hydrochloric and phosphorous acids (*vide* p. 350) to an aqueous or even acid solution of a weighed quantity of the mercurial compound, setting the mixture aside for twelve hours, collecting the precipitate on a tared filter, washing, drying at 100° C., and weighing (Rose). The experiment may be tried on half a gramme to a gramme of corrosive sublimate.

*Process* 3.—Two or three decigrammes of corrosive sublimate are dissolved in water, the solution acidulated with hydrochloric acid, excess of sulphuretted hydrogen passed through it, the precipitate collected on a tared filter, washed with cold water, dried at 100° C., and weighed.

*Proportional weights of equivalent quantities of mercury and its salts.*

| | | |
|---|---|---|
| Metal . . . . . | Hg . . . . | 200 |
| Mercurous chloride . . . | HgCl . . . . | 235.5 |
| Mercuric chloride . . . | $HgCl_2$ . . . . | 271 |
| Mercuric sulphide . . . | HgS . . . . | 232 |

## LEAD.

Lead is generally estimated either as (1) oxide, (2) sulphate, (3) chromate, or (4) metal.

*Process* 1.—Weigh out one or two grammes of pure acetate of lead in a covered crucible, previously tared, and heat slowly until no more vapors are evolved. Remove the lid, stir down the carbonaceous mass with a clean iron wire, and keep the crucible in the flame so long as any carbon remains unconsumed. Introduce some fragments of fused nitrate of ammonium, and again ignite until no metallic lead remains, and all excess of the nitrate has been decomposed. Cool and weigh the resulting oxide (PbO).

*Process* 2.—Dissolve 0.4 or 0.5 of a gramme of acetate of lead in a small quantity of water, drop in diluted sulphuric acid, add to the mixture twice its bulk of methylated spirit of wine, and set aside. Decant the supernatant liquid, collect the sulphate on a filter, wash with spirit, dry, transfer to a porcelain crucible, removing as much of the sulphate as possible from the paper, incinerate on the crucible-lid (not in the platinum coil, for the particles of reduced lead would unite with the platinum by fusion), ignite, cool, and weigh.

*Process* 3.—About half a gramme of acetate of lead is dissolved in two or three hundred c. c. of water, acetic acid added, and then solution of red chromate of potassium. Collect the precipitate on a tared filter, wash, dry at 100° C., and weigh.

*Process* 4.—In certain cases, notably in that of commercial white-lead, the lead may be estimated in the metallic state by means of cyanide of potassium. The lead paint (about 20 grammes) is weighed and carefully incinerated. The residue, a mixture of metallic lead and oxide of lead, is then mixed with several times its bulk of cyanide of potassium and the whole heated to fusion. With careful manipulation the lead collects in one globule, which, after cooling, may readily be separated from the mixed cyanide and cyanate and weighed. White lead, commercially pure, should contain 74 per cent. of lead.

*Molecular weight of salts of lead.*

| | | |
|---|---|---|
| Metal . . . . | Pb . . . . . | 207 |
| Acetate . . . . | $Pb2C_2H_3O_2,2H_2O$ . | 379 |
| Oxide . . . . | PbO . . . . . | 223 |
| Sulphate . . . | $PbSO_4$ . . . . . | 303 |
| Chromate . . . | $PbCrO_4$ . . . . . | 323.5 |

## SILVER.

Compounds of silver which are readily decomposed by heat are estimated in the form of (1) metal, others usually as (2) chloride (AgCl), but sometimes as (3) cyanide (AgNC).

*Process* 1.—Heat about a gramme of oxide of silver ($Ag_2O$) in a tared crucible, cool, and weigh. 232 of oxide yield 216 of metal. "29 grains heated to redness yield 27 grains of metallic silver."—*Brit. Pharm.*

*Process* 2.—Dissolve 0.4 or 0.5 of pure dry crystals of nitrate of silver in water, acidulate with two or three drops of nitric acid, slowly add hydrochloric acid, stirring rapidly, until no more precipitate falls. Pour off the supernatant liquid through a filter, wash the chloride of silver once or twice with hot water, transfer to the filter, complete the washing, and dry. After removing as much as possible of the precipitate from the paper to the crucible, burn the filter, letting its ash fall on the inverted lid of the crucible, moisten with a drop of nitric acid, warm, add a drop of hydrochloric acid, evaporate to dryness, replace the lid on the crucible, unite the whole until the edges of the mass of chloride begin to fuse; cool and weigh. 170 of nitrate yield 143.5 of chloride. According to the British Pharmacopœia, 10 parts of nitrate should thus yield 8.44 of chloride, and the filtrate from the chloride evaporated to dryness should leave no residue, indicating absence of nitrates of potassium or sodium and other similar adulterants.

*Process* 3.—Cyanide of silver may be collected on a tared filter and dried at 100° C. 170 of nitrate yield 134 of cyanide.

Silver and its salts may be volumetrically estimated by a standard solution of chloride of sodium.

*Cupellation.*—The amount of silver in an alloy may be also determined by a dry method. The metal is folded in a piece of thin sheet lead, placed on a *cupel* (*cupella*, little cup, made of compressed bone-earth) and heated in a furnace, the cupel being protected from the direct action of flame by a muff-shaped, or, rather, oven-shaped case termed a muffle. The metals melt, the baser become oxidized, the oxide of lead fusing and dissolving the other oxides; the fluid oxides

are absorbed by the porous cupel, a button of pure silver remaining. An alloy supposed to contain 95 per cent. of silver requires about 3 times its weight of lead for successful cupellation; if 92½ per cent. (English silver coin), between 5 and 6 times its weight of lead is necessary.

---

## QUESTIONS AND EXERCISES.

1035. Explain the gravimetric process by which the strength of the official solutions of ferric chloride, nitrate, and sulphate is determined.

1036. Mention the various amounts of ferrous and ferric salts equivalent to 100 parts of metal.

1037. State the precautions necessary to be observed in estimating arsenicum or antimony in the form of sulphide.

1038. In what form are the official compounds of bismuth weighed for quantitative purposes?

1039. Give an outline of the process by which mercury may be isolated from its official preparations and weighed in the metallic condition.

1040. Describe three methods for the quantitative analysis of salts of lead; and the weights of the respective precipitates, supposing 0.56 of crystallized acetate to have been operated on in each case.

1041. Describe the process by which silver is estimated in the forms of metal, chloride, and cyanide.

1042. What proportions of nitrate of silver are indicated, respectively, by 15 of metal, 9.8 of chloride, and 8.1 of cyanide?

1043. Describe cupellation.

---

# GRAVIMETRIC ESTIMATION OF THE ACIDULOUS RADICALS OF SALTS.

## CHLORIDES.

*Free chlorine* (chlorine water) and compounds which by action of acids yield free chlorine (Chlorinated Lime, Chlorinated Soda, and their official Solutions) are estimated volumetrically by a standard solution of hyposulphite of sodium (*vide* p. 565). The amount of combined chlorine in pure chlorides (HCl, NaCl) may also be determined by volumetric analysis with a standard solution of nitrate of silver (p. 556).

*Combined chlorine* is gravimetrically estimated in the form of chloride of silver, the operation being identical with that

just described for silver salts (p. 584). 58.5 parts of pure, colorless, crystallized chloride of sodium (rock-salt) yield 143.5 of chloride of silver.

## IODIDES.

*Free iodine* is estimated volumetrically by solutions of hyposulphite of sodium (*vide* p. 564).

*Combined iodine* is determined gravimetrically in the form of iodide of silver, the operations being conducted as with chloride of silver. Iodide of potassium may be used for an experimental determination: KI = 166 should yield AgI = 235. Of the official iodide of cadmium (*Cadmii Iodidum*, B. P.) it is stated that "ten grains dissolved in water, and nitrate of silver added in excess, give a precipitate which, when washed with water and afterwards with half an ounce of solution of ammonia, and dried, weighs 12.5 grains."

*In presence of chlorides and bromides* the iodine in iodides may be precipitated and weighed as iodide of palladium.

## BROMIDES.

*Free bromine* may be estimated by shaking with excess of solution of iodide of potassium, and then determining the equivalent quantity of liberated iodine by a standard solution of hyposulphite of sodium (p. 564).

*The bromine in bromides* may be precipitated and weighed as bromide of silver, the manipulations being the same as those for chloride of silver: 0.2 to 0.3 of pure bromide of potassium may be used for an experimental analysis.

## CYANIDES.

*The hydrogen cyanide* (hydrocyanic acid) is usually estimated volumetrically (*vide* p. 556).

From all soluble cyanides, cyanogen may be precipitated by nitrate of silver, after acidulating with nitric acid, the cyanide of silver collected on a tared filter, dried at 100° C., and weighed.

Of the official Diluted Hydrocyanic Acid, it is stated that one hundred grains (or 110 minims) precipitated by solution of nitrate of silver yield ten grains of dry cyanide of silver.

*Cyanide of Silver.*

| | | In 1 molecule. | In 100 parts. |
|---|---|---|---|
| Silver . . . | Ag . . | 107.93 . . | 80.59 |
| Cyanogen . | CN . . | 26.00 . . | 19.41 |
| | | 133.93 | 100.00 |

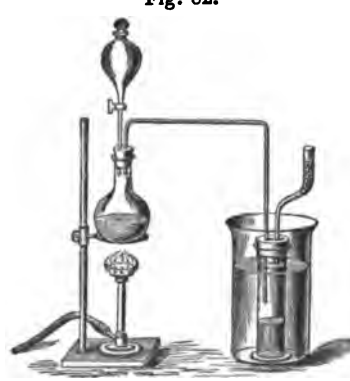
## NITRATES.

Nitrates cannot be estimated by direct gravimetric analysis, none of the basylous radicals yielding a definite nitrate insoluble in water. With some difficulty they may be determined by indirect volumetric methods.

*Process.*—The following (Thorpe's) method depends upon the fact (Gladstone and Tribe) that when zinc upon which copper is deposited in a spongy form is boiled with water, hydrogen is evolved. Thorpe found that in a solution containing nitrates the nascent hydrogen converts the whole of the nitrogen of the nitrates into ammonia, which may be collected and estimated. (The oxygen of the nitrate is simultaneously converted into water, the nitrate-metal into hydrate, and the zinc into hydrate of zinc. The power of the copper-zinc couple is considered to depend largely on the hydrogen absorbed by the finely divided metal.)

An apparatus such as shown in Fig. 82 should be constructed. A flask (about 100 c. c.) is fitted with an India-rubber cork, perforated for a delivery tube, which should be of strong glass tubing of about quarter-inch bore, and for a stoppered funnel, which should have about half the capacity of the flask. The whole is supported by a clamp or on wire-gauze. The outer jar shown in the figure should have a capacity of two or three litres, and the inner receiving-jar should be capable of holding 200 c. c. The latter is fitted with an India-rubber cork, perforated for the delivery-tube, and for another tube containing fragments of glass.

Fig. 82.



Estimation of Nitrates.

A few strips of clean zinc are boiled in a beaker with a two per cent. solution of sulphate of copper, the operation being repeated with a fresh portion of solution six consecutive times. A thick coating of finely divided copper is deposited. The pieces of metal are well washed and introduced into the flask, which is then half filled with pure water. To avoid transference, the flask itself may be used instead of the beaker. The funnel also is filled with pure water. Into the inner receiver is put a little pure water very slightly acidulated with hydrochloric acid; and the glass fragments are also moistened with the dilute acid (to prevent possible loss of ammonia). Water is now placed around the inner receiver in the outer jar, and, the connections being sound, heat is applied with the view of freeing the apparatus itself from any trace of ammonia. When the contents of the flask are evaporated nearly to dryness, pure water is admitted from the funnel until the flask is again about half full (the funnel should be filled again at once), and the distillation carried on as before. This must be repeated until no further trace of ammonia is evolved, when the apparatus is ready for use. On each occasion that the apparatus is used it must be freed from ammonia in this way. A suitable quantity of the substance to be estimated is now introduced (in the case of potable waters the prepared solid residue from 100 c. c., with an added fragment of recently ignited lime—the size of a hempseed—to promote the evolution of the ammonia), and water added, if necessary, until the flask is half full. Heat is now applied, and the operation conducted in the manner already described until ammonia ceases to come over—a point which always occurs in the case of water-residues when the flask has been refilled twice and the distillate is about 100 c. c. The warm water from the upper part of the cooling-jar may be removed by a siphon or otherwise, cold water being introduced from time to time.

The ammonia being all evolved, disconnect the flask and receiver simultaneously, and treat the contents of the latter by the Nessler method described on page 548. Urea yields but traces of ammonia by this process, and neither the sulphates nor chlorides of the alkali-metals effect the result.

*Another process* (Pelouze's improved by Fresenius) consists in adding the nitrate to an acid solution of a ferrous salt of known strength, and, when reaction is complete, estimating the amount of ferrous salt unattacked by volumetric solution of red chromate or of permanganate. Three molecular weights of converted ferrous salt indicate one molecular weight of nitric acid. Regeneration of nitric or nitrous acids by aerial oxidation of

the nitric oxide evolved is prevented either by a current of carbonic acid gas or by using a closed flask in which is a *Bunsen valve* (*i. e.*, a short attached piece of India-rubber tubing closed at the free extremity and having a sharp longitudinal slit in it a third of an inch long—a slit by which gases can escape, but cannot re-enter).

## SULPHIDES.

*Process* 1.—Soluble sulphides ($H_2S$, NaHS, *e. g.*) may be determined volumetrically by adding to the aqueous liquid a measured excess of an alkaline solution of arsenic of known strength, neutralizing by hydrochloric acid, diluting to any given volume, filtering off the sulphide of arsenicum precipitated, taking a portion of the filtrate equal to half or a third of the original volume, and, after neutralizing by acid carbonate of sodium, estimating the residual arsenic by the standard iodine solution (*vide* p. 558). The process may be tried on a measured volume of sulphuretted hydrogen (the weight of which is easily calculated; 1 litre of hydrogen = 0.0896 gramme) absorbed by a strong solution of soda or potash.

*Process* 2.—Sulphur and sulphides may also be quantitatively analyzed by oxidizing to sulphuric acid and precipitating in the form of sulphate of barium. A couple of decigrammes of a pure metallic sulphide may be decomposed by careful deflagration with a mixture of chlorate of potassium and carbonate of sodium, the product dissolved in water, acidulated with hydrochloric acid, solution of chloride of barium added, and the precipitated sulphate of barium purified and collected as described in connection with the estimation of barium (p. 573). Many sulphides may be oxidized in a flask by chlorate of potassium and hydrochloric acid, and then precipitated by chloride of barium. Experimental determinations may also be made on a weighed fragment of sulphur, about 0.1, cautiously fused with a solid caustic alkali, and the product oxidized while hot by the slow addition of powdered nitrate or chlorate of potassium, or, when cold, by treatment with chlorate of potassium and hydrochloric acid, and subsequent precipitation by chloride of barium.

*Note.*—Fusions performed by help of a gas-lamp must be carefully conducted; for any alkali that may creep over the side of a crucible will certainly absorb sulphurous acid from the products of combustion of the gas, and error result.

*Process* 3.—Soluble sulphides may also be treated with excess of an alkaline arsenite, arsenious sulphide be then preci-

pitated by the addition of hydrochloric acid, and the precipitate collected and weighed with the usual precautions (*vide* p. 579).

*Weights of equivalent quantities of sulphur and its compounds.*

| | | |
|---|---|---|
| Sulphur . . . . . | S . . . . . | 32 |
| Sulphuretted hydrogen . | $H_2S$ . . . . . | 34 |
| Sulphate of barium. . . | $BaSO_4$ . . . . . | 233 |
| Arsenious sulphide . . . | $(As_2S_3) \div 3$ . . . . | 82 |
| Bisulphide of iron . . . | $(FeS_2) \div 2$ . . . . | 60 |
| Sulphide of lead . . . | PbS . . . . . | 239 |

## SULPHITES.

Sulphites are usually estimated volumetrically by a standard solution of iodine (*vide* p. 558). Sulphites insoluble in water are diffused in that menstruum, hydrochloric acid added, and the iodine solution then dropped in.

If necessary, sulphites may be estimated gravimetrically by oxidation and precipitation in the form of sulphate of barium.

## SULPHATES.

These salts are always precipitated and weighed as sulphate of barium, the manipulations being identical with those performed in the determination of barium by means of sulphates (*vide* p. 573). The purity of Sulphate of Sodium (*Sodæ Sulphas*, B. P.), and the presence of not more than a given amount of sulphuric acid in Vinegar (*Acetum*, B. P.), are directed, in the British Pharmacopœia, to be ascertained by this process. Ten grains of sulphate of sodium yield 7.236 of sulphate of barium. Five ounces of vinegar should yield not more than about one-third of a gramme of sulphate of barium.

The amount of free sulphuric acid or hydrochloric acid in vinegar, lemon-juice, lime-juice, etc., may also be ascertained volumetrically by adding a known quantity of standard solution of soda, evaporating to dryness, incinerating, dissolving in water, and, by standard acid, estimating the quantity of soda still remaining free. The soda lost indicates the amount of free mineral acid (Hehner). Thresh estimates the chlorine in a sample of vinegar, adds a known additional amount of chlorine, preferably in the form of chloride of barium, evaporates, ignites; treats with water, adds bicarbonate of sodium, to remove excess of barium, filters, and again estimates the chlorine. A loss of 71 of chlorine ($Cl_2$) indicates 98 of free sulphuric acid ($H_2SO_4$).

The method of estimating free sulphuric, nitric, or hydrochloric acids proposed by Spence and Esilman is founded on their power of decolorizing a standard solution of ferric acetate.

*Proportional weights of equivalent quantities of sulphates.*

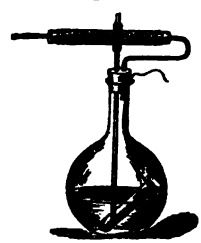
| | | |
|---|---|---|
| The sulphuric radical . . | $SO_4$ . . . . . | 96 |
| Sulphuric acid . . . . | $H_2SO_4$ . . . . . | 98 |
| Sulphate of barium . . . | $BaSO_4$ . . . . . | 233 |

## CARBONATES.

Carbonates are usually estimated by the loss in weight they undergo on the addition of a strong acid.

*Process* 1.—A small light flask is selected—of such a size that it can be conveniently weighed in a delicate balance. Two narrow glass tubes are fitted to the flask by a cork; the one straight, extending from about two or three centimetres above the cork to the bottom of the flask; the other cut off close to the cork on the inside and curved outwards so as to carry a thin drying-tube horizontally above the flask. (See Fig. 83.) The drying-tube is nearly filled with small pieces of chloride of calcium, a plug of cotton-wool preventing escape of any fragments at either end, and is attached by a pierced cord to the free extremity of the curved tube of the flask. A weighed quantity of any pure soluble carbonate is placed in the flask, a little water added, a miniature test-tube containing sulphuric acid lowered into the flask by a thread and supported so that the acid may not flow out, the cork inserted, the outer end of the piece of the straight glass tube closed by a fragment of cork or wax, and the whole weighed. The apparatus is then inclined so that the oil of vitriol and carbonate may slowly react; carbonic acid gas is evolved and escapes through the horizontal tube, any moisture being retained by the chloride of calcium. When effervescence has ceased, the gas still remaining in the vessel is sucked out; this is accomplished by adapting a piece of India-rubber tubing to the end of the drying-tube, removing the small plug from the straight tube, and aspirating slowly with the mouth for a few minutes. If the heat produced by the action of oil of vitriol and solution is considered insufficient to expel all the carbonic acid from the liquid, the plug is again inserted in the

Fig. 83.



Estimation of Carbonates.

tube and the contents of the flask gently boiled for some seconds. When the apparatus is cold, more air is again drawn through it, and the whole finally weighed. The loss is due to carbonic acid gas ($CO_2$), from the weight of which that of any carbonate is ascertained by calculation. Carbonates insoluble in water may be attacked by hydrochloric instead of sulphuric acid; granulated mixtures of carbonates and powdered tartaric or citric acids by inclosing the preparation in the inner tube and placing water in the flask, or *vice versâ*. The apparatus also may be modified in many ways to suit the requirements, convenience, or practice of the operator.

*Process* 2.—Carbonates from which carbonic acid gas is evolved by heat may be estimated by the loss they experience on ignition.

*Process* 3.—Free carbonic acid gas may be absorbed by a solid stick of potash or a strong alkaline solution, the loss in volume of the gas or mixture of gases indicating the amount originally present.

*Weights of equivalent quantities of carbonic acid gas and certain carbonates.*

| | | |
|---|---|---|
| Carbonic acid gas . . . . . | $CO_2$ . . . . . | 44 |
| Carbonic acid . . . . . | $H_2CO_3$ . . . . | 62 |
| Anhydrous carbonate of sodium . | $Na_2CO_3$ . . . . | 106 |
| Crystalline carbonate of sodium . | $Na_2CO_3,10H_2O$ . . | 286 |
| Anhydrous carbonate of potassium. | $K_2CO_3$ . . . . | 138 |
| Crystalline carbonate of potassium. | $K_2CO_3+16\%$aq. | 164.285 |
| Carbonate of calcium . . . . . | $CaCO_3$ . . . . | 100 |

## OXALATES.

*Process* 1.—The oxalic radical is usually precipitated in the form of oxalate of calcium, and weighed as carbonate, the manipulations being identical with those observed in the estimation of calcium (*vide* p. 574). The experiment may be performed on 0.3 or 0.4 of pure crystallized oxalic acid, 126 parts of which should yield 100 of carbonate of calcium.

*Process* 2.—Oxalates may also be determined by conversion of their acidulous radical into carbonic acid gas, and observation of the weight of the latter. The oxalate, water, and excess of black oxide of manganese are placed in the carbonic acid apparatus, page 591, a tube containing oil of vitriol lowered into the flask, the whole weighed, and the operation completed as for carbonate. From the following equation it will be seen

that every 88 parts of carbonic acid gas evolved indicate the presence of 126 parts of crystallized oxalic acid or an equivalent quantity of other oxalate :—

$$Na_2C_2O_4 + MnO_2 + 2H_2SO_4 = MnSO_4 + Na_2SO_4 + 2H_2O + 2CO_2.$$

The black oxide of manganese used in this experiment must be free from carbonates. The amount of materials employed is regulated by the size of the vessels.

## PHOSPHATES.

*Process* 1.—*From phosphates dissolved in water*, the phosphoric radical may be precipitated and weighed in the form of pyrophosphate of magnesium, the details of manipulation being similar to those observed in estimating magnesium (*vide* p. 575). Half a gramme or rather more of pure dry crystallized phosphate of sodium may be employed in experimental determinations. The official phosphate of ammonium (*Ammoniæ Phosphas*, B. P.) is quantitatively analyzed by this method. "If twenty grains of this salt be dissolved in water, and solution of ammonio-sulphate of magnesia added, a crystalline precipitate falls, which, when well washed upon a filter with solution of ammonia diluted with an equal volume of water, dried, and heated to redness, leaves 16.8 grains." Half a gramme or less is a more convenient quantity, if the operations be conducted with care. Solution of ammonio-sulphate of magnesium (B. P.) is prepared by dissolving 2 parts of sulphate of magnesium, 1 of chloride of ammonium, and 1 of solution of ammonia (20.6 per cent. $NH_4HO$) in 18 or 20 of distilled water; such a solution is of considerable use if several phosphoric determinations are about to be made.

*Process* 2.—*Free phosphoric acid* is most readily determined as phosphate of lead ($Pb_2PO_4$). Of the official solution of phosphoric acid it is stated that "355 grains by weight poured upon 180 grains of oxide of lead in fine powder leave, by evaporation, a residue (principally phosphate of lead) which, after it has been heated to dull redness, weighs 215.5 grains." One-tenth of these quantities may be used for experimental purposes; one to two grammes will give good results. The oxide of lead must be quite pure: it should be prepared by digesting red lead in warm dilute nitric acid, washing, drying, and heating a resulting puce-colored plumbic oxide in a covered porcelain crucible. The increase in weight obtained on evaporating a given amount of solution of phosphoric acid with a known

weight of perfectly pure oxide of lead (PbO) may be regarded as entirely due to phosphoric anhydride ($P_2O_5$).

$$3PbO + P_2O_5 = Pb_3(PO_4)_2,$$

the actual reaction being,

$$3PbO + 2H_3PO_4 = Pb_3(PO_4)_2 + 3H_2O.$$

From these equations, and the table of atomic weights (*vide* Appendix), the percentage of phosphoric acid ($H_3PO_4$) in any specimen of its solution may be easily calculated.

*Process* 3.—*The strength of pure solution of phosphoric acid* may be ascertained by specific gravity and reference to Tables.

*Process* 4.—Bone-earth, "superphosphate," the *Calcis Phosphas* of pharmacy, and other forms of phosphate of calcium known to be tolerably free from iron or aluminium, may be estimated by treating about half a gramme with hydrochloric acid somewhat diluted, filtering if necessary, warming, precipitating with excess of ammonia, collecting the precipitate ($Ca_3(PO_4)_2$), washing, drying, igniting, and weighing. "*Calcis phosphas*," if pure, will, in this process, lose no weight.

*Process* 5.—*Insoluble phosphates* in ashes, manures, etc., are treated as follows: a weighed quantity of the material (1.0 to 10.0) is digested in hydrochloric acid diluted with three or four times its bulk of water; filtered (insoluble matter and filter being thoroughly exhausted by water); ammonia added to the filtrate and washings until, after stirring, a faint cloudy precipitate is perceptible; solution of oxalic acid dropped in until, after agitation for a few minutes, the opalescence is destroyed; oxalate of ammonium next added, the whole warmed, oxalate of calcium removed by filtration, and the filtrate concentrated if very dilute; the liquid treated with citric acid in such quantity that ammonia when added in excess gives a clear *lemon-yellow* solution (Warington), magnesian mixture poured in (as in Process 1), and the precipitate of ammonio-magnesian phosphate collected, washed, dried, and weighed as already described in connection with the estimation of magnesium.

*Relative weights of equivalent quantities of phosphoric compounds.*

| | | |
|---|---|---|
| Phosphoric acid . . . . . | $H_3PO_4$ . . . . . | 98 |
| Pyrophosphate of magnesium . | ($Mg_2P_2O_7$=222) ÷2= | 111 |
| Phosphate of lead . . . . | ($Pb_3(PO_4)_2$=811) ÷2= | 405.5 |
| Phosphoric anhydride . . . | ($P_2O_5$=142) ÷2= | 71 |
| Phosphate of calcium . . . | ($Ca_3(PO_4)_2$=310) ÷2= | 155 |
| Superphosphate of calcium . | ($CaH_4(PO_4)_2$=234) ÷2= | 117 |

## QUESTIONS AND EXERCISES.

1044. What quantity of pure rock-salt is equivalent to 4.2 parts of chloride of silver?—*Ans.* 1.712.

1045. State the percentage of real iodide of potassium contained in a sample of which 8 parts yield 10.9 of iodide of silver?—*Ans.* 96.25.

1046. What is the strength of a solution of hydrocyanic acid 10 parts of which, by weight, yield .9 of cyanide of silver?—*Ans.* 1.81 per cent.

1047. How are nitrates quantitatively estimated?

1048. By what processes may the strength of sulphides be determined?

1049. How much real sulphate of sodium is contained in a specimen 10 parts of which yield 14.2 of sulphate of barium? — *Ans.* 86.34 per cent.

1050. Give details of the operations performed in the quantitative analysis of carbonates.

1051. What amount of carbonic acid gas should be obtained from 10 parts of acid carbonate (or bicarbonate) of potassium?—*Ans.* 4.4 parts.

1052. To what operation and what proportion of materials does the following equation refer?

$$Na_2C_2O_4 + MnO_2 + 2H_2SO_4 = MnSO_4 + Na_2SO_4 + 2H_2O + 2CO_2$$

1053. Explain the lead process for the estimation of phosphoric acid in the official solution.

1054. State the amount of superphosphate of calcium equivalent to 7.6 parts of pyrophosphate of magnesium?—*Ans.* 8.01.

---

## SILICATES.

Silica ($SiO_2$) may be separated from alkaline silicates, or from silicates decomposable by hydrochloric acid, by digesting the substance in hydrochloric acid at a temperature of 70° or 80° C. until completely disintegrated, evaporating to dryness, heating in an air-bath, again moistening with acid, diluting with hot water, filtering, washing, drying, igniting, and weighing.

## ESTIMATION OF WATER.

Water and other matters readily volatilized are most usually estimated by the loss in weight which a substance undergoes on being heated to a proper temperature. Thus, in the British Pharmacopœia, crystalline gallic acid ($H_3C_7H_3O_5,H_2O$) is stated to lose 9.5 per cent. of its weight at a temperature of 100° C.,

oxalate of cerium ($CeC_2O_4,3H_2O$) 52 per cent. on inceration, carbonate of potassium about 16 per cent. on exposure to a red heat, sulphate of quinine ($2C_{20}H_{24}N_2O_2,H_2SO_4,7H_2O$) 14.4 per cent. at 100° C., arseniate of sodium ($Na_2HAsO_4,7H_2O$) 40.38 per cent. at 149° C., carbonate of sodium ($Na_2CO_3,10H_2O$) 60.3 per cent., phosphate of sodium ($Na_2HPO_4,12H_2O$) 63 per cent., and sulphate of sodium ($Na_2SO_4,10H_2O$) 55.9 per cent. at a low red heat; oxide of bismuth heated to incipient redness should not diminish in weight.

*Process.*—One or two grammes of substance is sufficient in experiments on desiccation, the material being placed in a watch-glass, covered or uncovered porcelain crucible, or other vessel, according to the temperature to which it is to be exposed. Rapid desiccation at an exact temperature may be effected by introducing the substance into a tube having somewhat the shape of the letter U, sinking the lower part of the tube into a liquid kept at a definite temperature by aid of a thermometer, and drawing or forcing a current of dry air slowly through the apparatus. Substances liable to oxidation may be desiccated in a current of dried carbonic acid gas. The weights of the U-tube before and after the introduction of the salt, and after desiccation, give the amount of water sought. In all cases the material must be heated until it ceases to lose weight. Occasionally it is desirable to estimate water directly by conveying its vapor in a current of air through a weighed tube containing chloride of calcium and reweighing the tube at the close of the operation; the increase shows the amount of water.

*Note.*—Highly dried substances rapidly absorb moisture from the air; they must therefore be weighed quickly, inclosed, if possible, in tubes (p. 568), a pair of clamped watch-glasses, or a crucible having a tightly fitting lid.

## CARBON, HYDROGEN, OXYGEN, NITROGEN.

The quantitative analysis of animal and vegetable substances is either *proximate* or *ultimate*. Proximate quantitative analysis includes the estimation of water, oil, albumen, starch, cellulose, gum, resins, alkaloids, acids, glucosides, ash. It requires the application of much theoretical knowledge and manipulative skill, and cannot well be studied except under the guidance of a tutor. One of the best of the published works on the subject is by Rochleder, a translation of whose monograph will be found in the Pharmaceutical Journal, vol. i. 2d ser. pp. 562, 610; vol. ii. 2d ser. 24, 129, 160, 215, 274, 420, 478. Another is a small book, by Professor A. B. Prescott, "Outlines of Proximate Organic Analysis;" Van Nostrand, New York.

*Ultimate quantitative organic analysis* can only be successfully accomplished with the appliances of a well-appointed laboratory—a good balance, a gas-furnace giving a smokeless flame (7 or 8 centimetres wide and 70 or 80 centimetres long), special forms of glass apparatus, etc. *The theory of the operation* is simple: a weighed quantity of a substance is burnt to carbonic acid gas ($CO_2 = 44$) and water ($H_2O = 18$), and these products collected and weighed; 12 parts in every 44 of carbonic acid gas ($= \frac{3}{11}$) are carbon, 2 in every 18 of water ($= \frac{1}{9}$) are hydrogen; nitrogen if present escapes as gas. If nitrogen be a constituent, more of the substance is strongly heated with a mixture of the hydrates of sodium and calcium; these bodies then split up into oxides, oxygen, and hydrogen; the oxygen burns the carbon of the substance to carbonic acid gas, its hydrogen and nitrogen appearing as water and ammonia respectively; the carbonic acid and water are disregarded, the ammonia collected and weighed in the form of a double chloride of platinum and ammonium ($PtCl_4,2NH_4Cl = 447$), of which 28 parts in every 447 ($= \frac{1}{16}$) are nitrogen. The difference between the sum of the weights of hydrogen and carbon, and the weight of substance taken, is the proportion of oxygen in the body, supposing nitrogen to be absent. If nitrogen is present, the difference between the sum of the percentage of carbon, hydrogen, and nitrogen, and 100, is the percentage of oxygen. Shortly, carbon is estimated in the form of carbonic acid gas, hydrogen as water, nitrogen as ammonia, and oxygen by loss.

*The following is the outline of the necessary manipulation.* The source of the oxygen for the combustion of carbon and hydrogen is black oxide of copper in coarse powder. 200 or 300 grammes of this material are heated in a crucible to low redness for a short time to expel every trace of moisture—then transferred to store-tubes (Fig. 84) resembling test-tubes, half

Fig. 84.

a metre long, and having a slightly narrowed mouth, the tube being held in a cloth to protect the hand while the hot oxide is being directly introduced into the mouth of the tube by a scooping motion. As soon as the well-corked tube is cool, the oxide is poured, portion by portion, into a similar tube (the combustion-tube), but somewhat longer, drawn out to a quill (bent upwards nearly to a right angle) at one end, and not constricted at the mouth. Two such tubes are readily made by softening in the blowpipe-flame two or three centimetres of the central part of a tube about a metre long, and drawing the halves of the tube apart as shown in the following engraving (Fig. 85). The tubes are separated by melting the glass in the

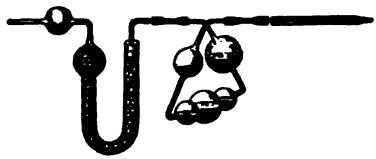
middle of the quilled portion. A few decigrammes of fused chlorate of potassium should first be dropped into the tube. After ten or fifteen centimetres of oxide have been poured in, about a decigramme of the substance to be analyzed is dropped

Fig. 85.

down the tube, then a few grammes of oxide, then another decigramme of substance, then more oxide, until three or four decigrammes of the body under examination have been added. The fifteen or twenty centimetres of alternate layers are next thoroughly mixed by a long copper wire having a short helix; more oxide is introduced, the wire cleansed by twisting the helix about in the pure oxide, and a plug of dry asbestos finally placed on the top of the oxide at about five centimetres from the mouth of the tube; the tube is then securely corked and set aside. The substance operated on may be pure white sugar, powdered, and dried; the tube in which it is contained is weighed before and after the removal of a portion for combustion; the loss is the quantity employed in the experiment. The combustion furnace may be such as shown on page 582. If the furnace is very powerful, or the combustion-tube not of the hardest glass, the tube should be inclosed in wire-gauze the elasticity of which has been destroyed by heating to redness. If the substance under experiment contains nitrogen, the plug of asbestos must be displaced by one of copper turnings, which serves to reduce any oxides of nitrogen, and thus insure the escape of nitrogen itself. The *water* produced when the prepared tube is heated, is collected in a small U-tube containing pieces of chloride of calcium, or pumicestone moistened with sulphuric acid (Fig. 86); the *carbonic acid gas* in a series of bulbs (Fig. 86) containing solution of potash (sp. gr. about 127). These bulbs may be purchased at any apparatus-shop. The chloride-of-calcium tube is fitted by a good cork to the combustion-tube, the potash bulbs by a short piece of India-rubber tubing to the chloride-of-calcium tube. The potash-bulbs may carry a short light tube containing a rod of caustic potash three or four centimetres long; this serves to arrest any moisture that might be carried away from the solution of pot-

ash by the dried expanded air which escapes during the operation. The combustion-tube having been placed in the furnace, and the drying-tube and potash-bulbs weighed and detached, the gas is lit under the asbestos, and, when the tube is red hot,

Fig. 86.



Chloride-of-calcium Tubes, and Potash-bulbs.

the flame slowly extended until nearly the whole tube is at the same temperature, the operation being conducted at such a rate that bubbles of gas escape through the potash-bulbs at about the rate of one per second. When no more gas passes, the extremity of the tube containing the chlorate of potassium is gently heated until oxygen ceases to be evolved; the quilled extremity of the combustion-tube is then broken, and air, dried and freed from carbonic acid, drawn slowly through the apparatus by suction through an India-rubber tube fixed on the free end of the potash-bulbs; perfect combustion of carbon and removal of all carbonic acid gas are thus insured. The drying-tube and bulbs are disconnected and weighed, the increase in weight due to carbonic acid gas and water respectively noted, and the percentages of carbon, hydrogen, and (by loss) oxygen calculated. This method is that of Liebig, with modification by Bunsen; good combustion-furnaces are those known as Hofmann's and Griffin's.

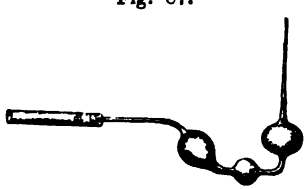
*Chromate of lead can be used for combustions in place of oxide of copper.* Its advantages are, its less hygroscopic nature, and the greater readiness with which it yields its oxygen to organic bodies when heated with them. It must not, however, be used with bodies containing nitrogen, since it would convert so large a proportion of the nitrogen into nitric oxide or higher oxide of nitrogen, that it would be necessary to use an inconveniently long layer of copper turnings to reduce these oxides and so prevent their absorption in the series of bulbs containing the solution of potash. Organic bodies, however, containing sulphur, bromine, iodine, or chlorine are burnt with advantage by means of chromate of lead. If oxide of copper were used with bodies containing sulphur, it would be necessary to place an additional tube containing per-

oxide of lead between the chloride-of-calcium tube and the potash-bulbs in order to absorb the sulphurous anhydride formed; this is entirely obviated by using chromate of lead, which itself retains the whole of the sulphur. Again, if bodies containing chlorine, iodine, or bromine are burnt by means of oxide of copper, then volatile chloride, iodide, or bromide of copper is formed, and, collecting in the chloride-of-calcium tube, vitiates the result with regard to the hydrogen; by using chromate of lead, however, the chlorine, iodine, and bromine are respectively retained in the combustion-tube as chloride, bromide, and iodide of lead.

In order to render the chromate fit for use, it is first fused and poured out on a clean iron plate; when cool it is powdered, and heated in a long tube throughout its whole length, while air, dried by passing through chloride of calcium or strong sulphuric acid, is drawn over it; when the color of the chromate changes to brown the heat can be withdrawn and the extremity of the tube furthest from the drying-apparatus closed, so that the air in passing into the tube on cooling may be quite dry; when cool the drying-tube is removed, the extremity securely corked, and the chromate of lead is ready for direct transference to the combustion-tube.

*The general manipulations for substances containing nitrogen* resemble the foregoing so far as the use of a combustion-tube and furnace and collection of the ammoniacal gas are concerned. The combustion-tube must be quilled at one end, and about a third of a metre long. The soda-lime is made by slaking quick-lime with a solution of soda, of such a strength that about two parts of quick-lime shall be mixed with one of hydrate of sodium, drying the product, heating to bright redness, and finely powdering; it should be preserved in a well-closed bottle. Some of the soda-lime is introduced into the tube, then layers of substance and soda-lime, mixture effected by a wire, more soda-lime added, and lastly a plug of asbestos. Bulbs (Fig. 87), known as those of Will and Varrentrapp (the originators of the method), containing hydrochloric acid of about 25 per cent., are then fitted by a cork, and the tube heated in a furnace. When gas ceases to pass, the quill is broken, and aspiration continued slowly until ammoniacal gas may be considered to have been all absorbed by the acid.

Fig. 87.



Nitrogen-bulbs.

The bulbs are disconnected, their contents and rinsings poured into a small dish, solution of perchloride of platinum added, and the operation completed as in the estimation of ammonium and potassium salts (*vide* pages 570 and 572).

*Liquids are analyzed* by a similar method to that adopted for solids, volatile liquids being inclosed in small bulbs having a long quill. These are weighed previously to and after the introduction of the liquid; just before being dropped into the combustion-tube the quill is broken.

*Formulæ.*—From the percentage composition of an organic substance an empirical formula may be deduced by dividing the weight of each constituent by its atomic weight, and converting the product into the simplest whole numbers; a rational formula by ascertaining the proportion in which the substance unites with a radical or body having a known combining proportion, etc. (*vide* p. 424).

*Chlorine*, *bromine*, or *iodine* contained in an organic substance is usually estimated by heating to redness a given weight of the material with ten times as much pure lime in a combustion-tube. Chloride, bromide, or iodide of calcium is thus produced. While still hot the tube is plunged into water, the mixture of broken glass and powder treated with pure diluted nitric acid in very slight excess; the filtered liquid precipitated by nitrate of silver, and the chloride, bromide, or iodide of silver collected, washed, dried, and weighed.

*Sulphur*, *phosphorus*, and *arsenicum* in organic salts may be estimated by gradually heating in a combustion-tube 1 part of the substance with a mixture of 10 parts nitre, 2 dried carbonate of sodium (in order to moderate deflagration), and 30 chloride of sodium. The product is dissolved in water acidulated by nitric acid, the sulphuric radical precipitated and estimated as sulphate of barium, the phosphoric and arsenic radicals as ammonio-magnesian phosphate or arseniate.

## QUINIA OR QUININE.

*De Vrij's Method for the Separation of the Mixed Alkaloids from Cinchona Bark.*—Twenty grammes of powdered and sifted bark, dried at 100° C., are mixed with milk of lime, made of 5 grammes of dry slaked lime and 50 grammes of water. This mixture is slowly dried, and, when entirely dry, heated in a flask with 200 c. c. of very strong alcohol (about 95 per cent.) till it boils. After well shaking, the clear liquid is poured by degrees into a percolating tube. The flask is then washed with 200 c. c. more of the same alcohol, and the liquid

passed through the percolator. After this has run through, a little clear colorless liquid may be further collected by pouring into the percolator successive small quantities of distilled water. The united clear liquors are now slightly acidulated with weak sulphuric acid, whereby a precipitate of sulphate of calcium is formed. When this has subsided, the greater part of the liquid may be decanted and the rest passed through a small filter. From the clear solution most of the spirit is distilled off, the residue poured into a capsule, and the still washed with a little weak spirit, the washings being added to the liquid in the capsule. The capsule is now heated on a water-bath till free from alcohol, and the remaining liquor, which contains all the alkaloids in the form of acid sulphates, is, after cooling, passed through a small filter. On the filter remains a mixture of quinovine* and fatty substances, which must be repeatedly washed with water slightly acidulated with sulphuric acid, till the washings are no longer rendered turbid by caustic soda. The filtrate, concentrated over a water-bath, is, while *still warm*, precipitated by a large excess of caustic soda. This precipitation from a warm solution has the advantage of rendering the alkaloids less voluminous, and therefore capable of being more easily washed. (Should the alkaloids melt—as is sometimes the case under this treatment—the liquid must be cooled and the fused alkaloids carefully powdered before proceeding to the next step of the process.) The precipitate is next placed on a small filter, washed with the smallest possible quantity of distilled water (sufficient to remove the soda salt without dissolving the quinine, etc.): the filter is laid on blotting-paper, and this is renewed until the mixed alkaloids can easily be removed from the filter without adhering to it,—this can be done before they are quite dry, but requires some practice. They are then heated in a tared capsule on a water-bath till they cease to lose weight. The observed weight multiplied by five gives the percentage of mixed alkaloids in the bark. The intensely yellow alkaline liquor separated from the alkaloids contains still a small quantity of quinovine which can be separated by acidifying with hydrochloric acid.

*De Vrij's Method for the Separation and Quantitative Determination of all the different Cinchona Alkaloids.*—This is based upon the following facts (nomenclature adopted by the Quinological Congress of Amsterdam, 1877):—

---

* This is strictly a mixture of quinovine and quinovic acid, but, as quinovine is by far the larger constituent, I actually prefer the name of quinovine for the mixture.—*De Vrij.*

1. The great solubility of quinine and amorphous alkaloids in ether, and the *relative* insolubility of quinidine, cinchonidine, and cinchonine in this liquid.

2. The great solubility of the iodosulphate of the amorphous alkaloid in alcohol of at least 90 per cent., and the very slight solubility of the iodosulphate of quinine (herapathite) in the same.

3. The great difference in solubility between the tartrate of cinchonidine and the tartrates of cinchonine and quinidine—the first being soluble in 1265 parts of water at 10° C., the second in 35.6 parts of water at 16° C., and the third in 38.8 parts of water at 15° C.

4. The great difference in solubility between the hydriodate of quinidine and the hydriodates of cinchonidine and cinchonine in water and alcohol. One part of hydriodate of quinidine requires 1250 parts of water at 15° C. or 110 parts of alcohol; the same salt of cinchonidine 110 parts of water or 3 parts of alcohol; and that of cinchonine 128 of water or 3 of alcohol.

These facts are applied to the separation and determination of the different cinchona alkaloids in the following manner:—

A. *Determination of Quinine.*—Two grammes of the pulverized mixed alkaloids are mixed with 30 grammes of ether, and the mixture, after well shaking, left at rest till the next day. By this operation all the quinine and amorphous alkaloid are dissolved, together with traces of quinine and cinchonidine, while the chief part of these, together with cinchonine, remains undissolved.

After filtering the ethereal solution and washing the residue with ether, the united filtrates are distilled and the weight of the residue ascertained. This is dissolved in 40 times its weight of alcohol of 90 or 92 per cent., containing .7562 per cent. of sulphuric acid, to obtain an alcoholic solution of the acid sulphates of the alkaloids. From this solution the quinine is separated by adding carefully, by means of a pipette, an alcoholic solution of *iodosulphate of chinioidine*,* as long as

* This reagent is thus prepared: Two parts of *Sulphate of Chinioidine* are dissolved in 8 parts of water, containing 5 per cent. of sulphuric acid. (The salt is identical with *Sulphate of Amorphous Quinine, prepared according to Dr. de Vrij's process*, by Messrs. Howard and Sons, Stratford.) To this *clear* solution, contained in a large capsule, a solution of one part of iodine and two parts of iodide of potassium in 100 parts of water is *slowly* added under continuous stirring so that no part of the solution of chinioidine comes into contact with an excess of iodine. By this addition an orange-colored flocculent precipitate is formed of iodosulphate of chinioidine, which either spontaneously or

a dark brown or red precipitate of iodosulphate of quinine (herapathite) falls. As soon as all the quinine has been precipitated and a slight excess of the reagent added, the liquid acquires an intense yellow color. The beaker containing the liquor and precipitate is now covered with a watch-glass and heated on a water-bath till the liquid *begins* to boil. After cooling the beaker is weighed to ascertain the amount of liquid present; the latter is then poured on to a filter placed in a tared funnel, and the precipitate washed with a saturated solution of herapathite in alcohol (92 per cent.). The weight of the moist filter is now taken, and the precipitate allowed to dry in the funnel, and its weight again taken. It is then removed from the funnel to a watch-glass and completely dried over a water-bath; another closely fitting glass is then placed over it and its weight is once more taken, the weighing being repeated till constant. The *rationale* of the latter part of the process is as follows: Herapathite is soluble in alcohol at 15° C. to the extent of .125 per cent. The quantity of herapathite held in solution by the alcohol present after the addition of the precipitant used is to be calculated and added to the final amount obtained. The difference in weight between the moist and the dry filter (weighed in the funnel) gives the amount of alcohol driven off in drying, and from this can be calculated the amount of herapathite left on the filter from the saturated solution employed in the washing. A final correction can then be made by subtracting this from the sum obtained as above—the tem-

---

by a slight elevation of temperature collapses into a dark brown-red colored resinous substance, whilst the supernatant liquor becomes clear and slightly yellow-colored. This liquor (still containing chinioidine, which can be obtained in a very pure state, if a little sulphurous acid is added before precipitating the alkaloid by caustic soda) is poured off, and the resinous substance is washed by heating it on a water-bath with distilled water. After washing, the resinous substance is heated on the water-bath till all the water has been evaporated. It is then soft and tenacious at the temperature of the water-bath, but becomes hard and brittle after cooling. One part of this substance is now heated with 6 parts of alcohol of 92 or 94 per cent. on a water-bath, and is thus dissolved, and the solution allowed to cool. In cooling, a part of the dissolved substance is separated. The clear dark-colored solution is evaporated on a water-bath, and the residue dissolved in 6 parts of cold alcohol. This second solution leaves a small part of insoluble substance. The clear dark-colored solution obtained by the separation of this insoluble matter, either by decantation or filtration, constitutes the reagent which De Vrij, under the name of *iodosulphate of chinioidine*, uses both for the qualitative and quantitative determination of *crystallizable* quinine.

perature of the laboratory being taken into account in both cases. The resulting correction may be thus expressed:—

$$\left.\begin{matrix}\text{Total weight}\\ \text{of herapathite}\end{matrix}\right\} = \text{wt. obtained} + .00125 \times \left\{\begin{matrix}\text{total weight of alcohol present} - \\ \text{loss of alcohol in drying precip.}\end{matrix}\right.$$

From this weight the amount of *crystallizable* quinine can be calculated by the use of Jörgensen's formula—$4C_{20}H_{24}N_2O_2$, $3H_2SO_4$,$2HI$,$I_4$=2354; according to which 1 part of iodosulphate of quinine, dried at 100° C., represents .55055 of anhydrous quinine, or .734 of pure commercial disulphate of quinine.

B. *Determination of the other Alkaloids.*—Two grammes of the pulverized mixed alkaloids are dissolved in weak hydrochloric acid to obtain a slightly alkaline solution measuring 70 c. c. By adding 1 gramme of Rochelle salt to this solution the tartrates of quinine and cinchonidine are separated; these are collected on a filter, washed with a little water, and dried on a water-bath. One part of these tartrates represents .80844 of quinine and cinchonidine: from the amount of these alkaloids thus found the amount of quinine already ascertained is subtracted, the remainder representing the cinchonidine present. In the filtrate from the tartrates quinidine (if present) is precipitated by a concentrated solution of potassium iodide; one part of the dried hydriodate representing .86504 of crystallized quinidine. The remaining solution is treated with caustic soda, and the precipitate (if any) washed with ether. The residue represents the amount of cinchonine. Finally, by distilling the ether from the washings can be ascertained the amount of amorphous alkaloid, which often, in the case of analysis of Indian barks, contains traces of quinamine.

*A quantitative determination of the purity of commercial sulphate of quinine* may be made by dissolving 1 gramme of the salt in 20 grammes of alcohol containing .7562 per cent. of sulphuric acid, adding 20 more grammes of alcohol, and precipitating the quinine as herapathite in the manner above described. If pure, one gramme of quinine should yield 1.3 gramme of herapathite.

*Paul's Modification of Kerner's Test for Sulphate of Cinchonidine in Sulphate of Quinine.*—Dissolve 4 or 5 grammes of the salt in 80 to 150 c. c. of boiling water. Set aside for some hours, or during a night. Most of the quinine sulphate is deposited, while the more soluble cinchonidine sulphate remains in solution. Filter. To the filtrate in a bottle add ether,

shaking occasionally, until a distinct layer of ether remains undissolved. Add now ammonia in excess; quinine is precipitated and then redissolved by the excess of ammonia, while the cinchonidine remains insoluble. Set this mixture aside for a few hours; cinchonidine, which may have been carried into solution in the ether along with the quinine, will separate out in a crystalline state. If a weighed quantity of the original salt has been operated on, the separated cinchonidine may be collected on a filter, washed with a little ether, and weighed. Care must be taken that the filter does not retain quinine deposited by the evaporating of the ether which held it in solution. This will be avoided if the flask in which the operation is conducted is of such a size that the ethereal solution of quinine collects in the neck; for the latter liquid may be removed by a pipette, a little more ether then poured in, the cork replaced, and the whole shaken and set aside; and this operation repeated two or three times before the cinchonidine is collected. The cinchonidine being thus washed with a minimum of ether suffers a minimum loss.

The quinine sulphate deposited from the first aqueous solution should be again recrystallized, and its mother-liquor treated as above. And even the sulphate deposited in this second treatment should be once more carried through the operation. The whole of the cinchonidine may finally be collected on one filter.

The aqueous filtrates may even be evaporated, set aside, filtered, and the ether test again applied (Parker).

Sulphate of quinine normally contains 14.45 per cent. of water; sulphate of cinchonidine 13.17 per cent.; all given off at 100° C. The drying should therefore be effected at 100° C., and the dried salt weighed in well-fitting weighing tubes. 100 parts of cinchonidine are equivalent to 116 parts of sulphate of cinchonidine.

*Of the Citrate of Iron and Quinia* (*Ferri et Quiniæ Citras*, B. P. and U. S. P.) it is officially stated that "fifty grains dissolved in a fluidounce of water and treated with a slight excess of ammonia give a white precipitate which, when collected on a filter, washed, and dried (at 260° F.), weighs eight grains. The precipitate is almost entirely soluble in two or three fluidrachms of pure ether," and the ethereal solution set aside for twelve hours in a small well-corked bottle yields no crystalline deposit (of quinidia).

## MORPHIA.

*The official process for the estimation of this alkaloid in opium* is conducted in the following manner:—

Take of opium 100 grains, slaked lime 100 grains, distilled water 4 ounces. Break down the opium, and steep it in an ounce of the water for twenty-four hours, stirring the mixture frequently. Transfer it to a displacement-apparatus, and pour on the remainder of the water in successive portions, so as to exhaust the opium by percolation. To the infusion thus obtained, placed in a flask, add the lime, boil for ten minutes, filter, wash the residue on the filter with an ounce of boiling water. Acidulate the mixed filtrates slightly with diluted hydrochloric acid, evaporate it to the bulk of half an ounce, and let it cool. Neutralize cautiously with solution of ammonia, carefully avoiding an excess; remove by filtration the brown matter which separates, wash it with an ounce of hot water, mix the washings with the filtrate, concentrate the whole to the bulk of half an ounce, and add now solution of ammonia in slight excess. After twenty-four hours collect the precipitated morphia on a weighed filter, wash it with cold water, and dry it at 212° F. It ought to weigh at least from six to eight grains.

*Of Hydrochlorate of Morphia* it is stated that "twenty grains of the salt, dissolved in half an ounce of warm water, with ammonia added in the slightest possible excess, give on cooling a crystalline precipitate which, when washed with a little cold water and dried by exposure to the air, weighs 15.18 grains."

*Testing Opium.*—Professor Schneider has proposed, in the 6th revised edition of the Pharmacopœia Austriaca, the following method for testing the quality of opium. Ten grammes of previously dried and powdered opium are treated with a mixture of 150 grammes of distilled water, to which 20 grammes of pure hydrochloric acid, sp. gr. 1.12, are added; the residue, after extraction, should not exceed 4.5 grammes weight; to the acid fluid 20 grammes of common salt are added, and the precipitate thereby caused is collected, after twenty-four hours, on a filter, and the latter washed with a solution of common salt; to the filtrate ammonia is added, and the fluid left standing again for twenty-four hours; the crystals which have separated are collected, redissolved in acetic acid, and precipitated with ammonia; the precipitate so obtained is washed, dried, and weighed; its weight should not be less than 1 gramme.

Flückiger and Hanbury also recommend a good process ("Pharmacographia," p. 59).

## SUGAR.

The qualitative test of sugar, by means of an alkaline copper solution (*vide* p. 409), may be applied in the estimation of sugar in sacchariferous substances.

*Process.*—34.65 grammes of pure dry crystals of ordinary sulphate of copper are dissolved in about 250 c. c. of distilled water, and 173 grammes of pure crystals of the double tartrate of potassium and sodium are dissolved in 480 c. c. of solution of caustic soda of sp. gr. 1.14. The solutions are mixed and water added until one litre of solution is obtained. 100 c. c. of this solution represent 3.464 grammes of sulphate of copper, and correspond to 0.5 of a gramme of pure anhydrous grape-sugar, 0.475 of cane-sugar, or 0.45 starch. It must be preserved in a well-stopped bottle to prevent absorption of carbonic acid, and be kept in a dark place. If it give a precipitate on boiling, a little solution of soda may be added in making experiments.

Dissolve 0.472 of pure dry powdered cane-sugar in about 50 c. c. of water, convert into grape-sugar by acidulating with sulphuric acid and boiling, neutralize with carbonate of sodium, and dilute to 100 c. c. Place 10 c. c. of the copper solution in a small flask, dilute with three or four times its bulk of water and gently boil. Into the boiling liquid drop the solution of sugar from a burette, one cubic cintimetre, or less, at a time, until, after standing for the precipitate to subside, the supernatant liquid has just lost its blue color; 10 c. c. of the solution of the sugar should be required to produce this effect, =0.0475 of cane-sugar or 0.05 of grape-sugar. Experiments on pure cane-sugar must be practised until accuracy is attained; syrups, diabetic urine, and saccharated substances containing unknown quantities of sugar may then be analyzed.

*Starch* is converted into grape-sugar by gentle ebullition with dilute acid for eight or ten hours, the solution being finally diluted so that one part of starch, or rather sugar, shall be contained in about 150 of water.

*Saccharimetry.*—A generic term for certain volumetric operations undertaken with the view of ascertaining the quantity of sugar present in any matter in which it may be contained.

Saccharimetry is frequently performed upon common syrup (*Syrupus*, B. P.) and solutions which are known to contain nothing but cane- (ordinary) sugar, the object being merely to ascertain the amount present. In such a case it is only necessary to take the specific gravity of the liquid at 60° F., and

then refer to a previously prepared Table of density and percentages.

| Specific gravity. | Sugar per cent. | Specific gravity. | Sugar per cent. | Specific gravity. | Sugar per cent. |
|---|---|---|---|---|---|
| 1.007 . . | 1.8 | 1.100 . . | 23.7 | 1.210 . . | 46.2 |
| 1.014 . . | 3.5 | 1.108 . . | 25.6 | 1.221 . . | 48.1 |
| 1.022 . . | 5.2 | 1.116 . . | 27.6 | 1.231 . . | 50.0 |
| 1.029 . . | 7.0 | 1.125 . . | 29.4 | 1.242 . . | 52.1 |
| 1.036 . . | 8.4 | 1.134 . . | 31.5 | 1.252 . . | 54.1 |
| 1.044 . . | 10.4 | 1.143 . . | 33.4 | 1.261 . . | 56.0 |
| 1.052 . . | 12.4 | 1.152 . . | 35.2 | 1.275 . . | 58.0 |
| 1.060 . . | 14.4 | 1.161 . . | 37.0 | 1.286 . . | 60.1 |
| 1.067 . . | 16 3 | 1.171 . . | 38.8 | 1.289 . . | 62.2 |
| 1.075 . . | 18.2 | 1.180 . . | 40.6 | 1.309 . . | 64.4 |
| 1.083 . . | 20.0 | 1.190 . . | 42.4 | 1.321 . . | 66.6? |
| 1.091 . . | 21.8 | 1.199 . . | 44.3 | 1.330 (B. P.) | 66.6? |

The sp. gr. may be taken by an hydrometer, technically termed a *saccharometer*. (The above spec. gravs. = 1° to 35° Baumé.)

If a liquid contains other substances besides cane-sugar, the test of specific gravity is of little or no value. Advantage may then be taken of the fact that syrup causes right-handed twisting of a ray of polarized light to an extent exactly proportionate to the amount of sugar in solution. The saccharine fluid is placed in a long tube having opaque sides and transparent ends; and a ray of homogeneous light, polarized by reflection from a black-glass mirror or otherwise, is sent through the liquid and optically examined by a plate of tourmaline, Nicol's prism, or other polarizing eyepiece. Attached to the eyepiece is a short arm which traverses a circle divided into degrees. The eyepiece and arm are previously so adjusted that when the ray is no longer visible the arm points to the zero of the scale of degrees. The saccharine solution, however, so twists the ray as to again render it visible; and the number of degrees which the eyepiece has to be rotated before the ray is once more invisible is exactly proportionate to the strength of the solution. The value of the degrees having been ascertained by direct experiment and the results tabulated, a reference to the Table at once indicates the percentage of sugar in the liquid under examination. Grape-sugar also possesses the property of dextro-rotation, but less powerfully than cane-sugar; moreover the former variety does not, like cane-sugar, suffer inversion of the direction of rotation on the addition of hydrochloric acid to its solution—an operation that furnishes data for ascer-

taining the amounts of cane- and of grape-sugar, or of crystallizable and non-crystallizable sugar, present in a mixture. In using the polariscope-saccharometer, it is convenient to employ tubes of uniform size, and always to operate at the same temperature. Various modes are adopted of applying, for the purposes of quantitative analysis, the action of syrup on polarized light.

## ALCOHOL.

*Mulder's process* for the determination of the amount of alcohol in wines, beer, tinctures, and other alcoholic liquids containing vegetable matter is as follows: Take the specific gravity and temperature of the liquid, and measure off a certain quantity (100 cubic centimetres); evaporate to one-half or less, avoiding ebullition in order that particles of the material may not be carried away by the steam. Dilute with water to the original bulk, and take the specific gravity at the same temperature as before. Of the figures representing this latter specific gravity, all over 1.000 show to what extent dissolved solid matter affected the original specific gravity of the liquid. Thus, the specific gravity of a sample of wine at 15°.5 C. is 0.9951; evaporated till all alcohol is removed and diluted with water to the original bulk, the specific gravity at 15°.5 C. is 1.0081; 0.0081 represents the gravitating effect of dissolved solid matter in 0.9951 parts of original wine. 0.0081 subtracted from 0.9951 leaves 0.987, which is the specific gravity of the water and alcohol of the wine. On referring to a Table of the strengths of diluted alcohol of different specific gravities (p. 634), 0.987 at 15°.5 C. is found to indicate a spirit containing 8 per cent. of real alcohol. If the foregoing operation be conducted in a retort, the liquid being boiled and the steam carefully condensed, the distillate, diluted with water to the original bulk of wine operated on, will still more accurately represent the amount of water and alcohol in the wine—its specific gravity showing the percentage of real alcohol present.

---

## DIALYSIS.

Dialysis (from διά, *dia*, through, and λύσις, *lusis*, a loosing or resolving) is a term applied by Graham to a process of analysis by diffusion through a septum. The apparatus used in the process is called a *dialyzer*, and is constructed and employed in the following manner. The most convenient septum is the commercial article known as *parchment paper*, made by

immersing unsized paper for a short time in sulphuric acid; it is sold by most dealers in chemical apparatus. A piece of this material is stretched over a gutta-percha hoop, and secured by a second external hoop. Dialyzers of useful size are one or two inches deep and five to ten inches wide. Liquids to be dialyzed are poured into the dialyzer, which is then floated in a flat dish containing distilled water. The portion passing through the septum is termed the *diffusate*, the portion which does not pass through is termed the *dialysate*.

The practical value of dialysis depends upon the fact that certain substances will diffuse through a given septum far more rapidly than others. Uncrystallizable bodies diffuse very slowly. Of such matters as starch, gum, albumen, and gelatine, the last named is perhaps least diffusive; hence substances of this class are termed *colloids*, or bodies like *collin*, which is the soluble form of gelatine. Substances which diffuse rapidly are most crystalline; hence bodies of this class are termed *crystalloids*.

Solution of two parts of the following named substances in 100 parts of distilled water were dialyzed by Graham for twenty-four hours. The amounts of each substance which passed through the septum bore the following relations to one another:—

| | |
|---|---|
| Chloride of sodium | 1000 |
| Ammonium | 847 |
| Theine | 703 |
| Salicin | 503 |
| Cane-sugar | 472 |
| Amygdalin | 311 |
| Extract of logwood | 168 |
| Catechu | 159 |
| Extract of cochineal | 51 |
| Gallo-tannic acid | 30 |
| Extract of litmus | 19 |
| Purified caramel | 5 |

Ten-per-cent. solutions, under similar circumstances, gave the following results:—

| | |
|---|---|
| Gum Arabic | 4 |
| Starch-sugar | 266 |
| Cane-sugar | 214 |
| Glycerine | 440 |
| Alcohol | 476 |
| Chloride of sodium | 1000 |

*Dialyzed iron*, an aqueous solution of about 5 per cent. of highly basic oxychloride of iron, is obtained by saturating solution of perchloride of iron with ferric hydrate, by adding ammonia, or, better, carbonate of sodium, and shaking vigorously until the precipitated hydrate ceases to redissolve, filtering if necessary, placing on a dialyzer floating in distilled water, and displacing the fluid in the dish by water daily for a week or two, or until the diffusate gives no reaction with nitrate of silver. The crystalloids (chloride of sodium or other salt) pass through the dialyzer; the colloid fluid which does not pass through the dialyzer is the highly basic oxychloride of iron, or so-called "dialyzed iron," or "dialytic iron."

The phenomena of dialysis show that crystalloids are superior to colloids in affinity for water. If a solution of chloride of sodium be placed at the bottom of a jar, and covered by a hot solution of gelatine of sufficient strength to solidify on cooling, the chloride of sodium will diffuse up into the solid jelly, because the water of the solid jelly has a greater affinity for the salt than it has for the gelatine. The solid jelly may obviously be reduced in thickness, and saline liquids placed above it; indeed the conditions would then be still more favorable for diffusion. Replace the stratum of jelly by a permanent colloid, such as parchment paper; the result is the same, but the permanent character of the septum admits of its practical application.

Further researches on dialysis will probably throw much light on several important points in connection with physiological chemistry; for there is little doubt that alimentary matter passes through the cell-walls of animals and plants by this process.

---

## QUESTIONS AND EXERCISES.

1055. Carbonate of potassium is said to lose 16 per cent. of water, on exposure to a red heat; give the details of manipulation observed in verifying this statement.

1056. Write a few paragraphs descriptive of the process of ultimate organic analysis.

1057. In what forms are carbon, hydrogen, and nitrogen weighed in quantitative analysis?

1058. In the combustion of .41 of a gramme of sugar, what weights of products will be obtained?—*Ans.* .632 of carbonic acid gas ($CO_2$) and .237 of water ($H_2O$).

1059. How is cinchona assayed for mixed alkaloids?

1060. On what facts does De Vrij found his method for the separation and quantitative determination of all the cinchona alkaloids?

1061. Describe De Vrij's process for the assay of commercial sulphate of quinia.

1062. Give the official method for the estimation of morphia in opium.

1063. Mention the operations necessary for the estimation of the proportion of sugar in saccharated carbonate of iron, or in a specimen of diabetic urine.

1064. What is understood by *Saccharimetry?*

1065. Give two processes for the estimation of the percentage of alcohol in tinctures, wines, or beer.

1066. Define dialysis.

---

## CONCLUSION.

Detailed instructions for the quantitative analysis of potable water, articles of food, general technical products, special minerals, soils, manures, air, illuminating agents (including solid fats, oils, spirits, petroleum, and gas), dyes, and tanning-materials, would scarcely be in place in this volume.

The course through which the reader has been conducted will, it is hoped, have taught the principles of the science of chemistry, and given special knowledge concerning the applications of that science to medicine and pharmacy, as well as have imparted sufficient manipulative skill to meet the requirements of manufacture or analysis. The author would venture to suggest that this knowledge be utilized, not only in the way of personal advantage, but in experimental researches on chemical subjects connected with therapeutics and pharmacy. The discovery and publication of a new truth, great or small, is the best means whereby to aid in advancing the calling in which we may be engaged, benefit our fellow-creatures and ourselves, and unveil the laws of the Creator of all things.

# APPENDIX.

## TABLE OF OFFICIAL TESTS FOR IMPURITIES IN (B. P.) PHARMACOPŒIAL PREPARATIONS.*

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Acacia Gummi | Starch | Iodine | 275 |
| Acetum | More than one thousandth of Sulphuric acid | Quantitative analysis | 590 |
| Acidum Aceticum | Traces of Lead or Copper | Sulphuretted Hydrogen | 222 |
| | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Hydrochloric acid | Nitrate of Silver | 269 |
| | Sulphurous acid | Nascent Hydrogen | 307 |
| | Mineral matter | Evaporation and ignition | 101 |
| Acid. Acetic. Glaciale | Sulphurous acid | Nascent Hydrogen | 307 |
| Acidum Boracicum | Alkaline salts | Insolubility in Alcohol | 333 |
| Acidum Carbolicum | Creasote | Oxidation, etc. See Creasotum | 444 |
| Acidum Citricum | Traces of Copper or Lead | Sulphuretted Hydrogen | 222 |
| | Tartaric acid | Acetate of Potassium | 322 |
| | Tartaric acid | Excess of Lime-water | 326 |
| | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Mineral matter | Incineration | 101 |
| Acidum Gallicum | Tannic acid | (Isinglass) Gelatine | 356 |
| Acidum Hydrochloricum | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Arsenic | Sulphuretted Hydrogen, Copper | 172 |
| | Sulphurous acid | Nascent Hydrogen | 307 |
| Acidum Hydrocyanicum Dilutum | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Hydrochloric acid | Ppt. by Nitr. silver, insol. in nitric acid | 269 |
| Acidum Nitricum | Mineral matter | Evaporation and ignition | 101 |
| | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Hydrochloric acid | Nitrate of Silver | 269 |
| | Mineral matter | Incineration | 99 |
| Acidum Oxalicum | Lead or Platinum | Sulphuretted Hydrogen | 258 |
| Acidum Phosphoricum Dilutum | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| | Hydrochloric acid | Nitrate of silver and Nitric acid | 269 |
| | Metaphosphoric acid | Albumen | 348 |
| | Nitric acid | Ferrous Sulphate and Sulphuric acid | 289 |
| | Phosphorous acid | Corrosive sublimate | 350 |
| | Ammonium (phosphate) | Hot alkali | 98 |

* The manipulations necessary to be observed in testing for impurities will be found described in the paragraphs treating of those substances. The Table also includes references to processes for ascertaining deficiency in strength of official articles.

The other "characters and tests" of pharmacopœial compounds have been given in connection with the respective synthetical and analytical reactions.

| Name of Preparation. | Impurities. | Tests. | Page |
|---|---|---|---|
| Acidum Sulphuricum | Mineral matter | Evaporation and ignition | 101 |
| | Nitric acid | Ferrous Sulphate | 289 |
| | Arsenic or Lead | Sulphuretted Hydrogen | 223 |
| Acidum Sulphurosum | Sulphuric acid | Chloride or Nitrate of Barium | 311 |
| Acidum Tannicum | Mineral matter | Incineration | 101 |
| Acidum Tartaricum | Metallic matter, as Lead | Sulphuretted Hydrogen | 222 |
| | Oxalic acid | Sulphate of Calcium | 317 |
| | Calcium (tartrate or sulphate). | Oxalate of Ammonium | 115 |
| Aconitia | Mineral matter | Incineration | 101 |
| Adeps præparatus | Mineral matter | Incineration | 101 |
| | Chloride (of sodium) | Nitrate of Silver | 269 |
| | Starch (flour) | Iodine | 275 |
| Æther | Alcohol | Boiling point, sp. gravity | 429 |
| Æther Aceticus | Water | Specific gravity | 431 |
| | Alcohol | Boiling point | |
| Æther Purus | Alcohol, Water | Specific gravity | 429 |
| Alcohol | Resin or oil | Opalescence on dilution | 425 |
| | Water | Anhydrous Sulphate of Copper | 192 |
| Alcohol Amylicum | Other spirituous matter | Boiling point, sp. gravity | 441 |
| Alum | Iron (sulphate) | Yellow or red prussiate | 160 |
| Ammonii Benzoas | Fixed salts | Non-volatility | 101 |
| Ammonii Bromidum | Free Bromine | Odor | 270 |
| | Iodide (of Ammonium) | Chlorine water and starch | 275 |
| | General | Quantitative analysis | 564 |
| Ammonii Carbonas | Fixed salts | Non-volatility | 101 |
| | Sulphate (of Ammonium) | Chloride or Nitrate of Barium | 311 |
| | Chloride (of Ammonium) | Nitrate of Silver | 269 |
| Ammonii Nitras | Chloride of Ammonium | Nitrate of Silver | 269 |
| | Sulphate of Ammonium | Chlor. or Nit. of Barium | 311 |
| Ammonii Chloridum | Fixed salts | Non-volatility | 101 |
| Ammonii Nitras | Sulphate (of Ammonium) | Chloride of Barium | 311 |
| | Chloride (of Ammonium) | Nitrate of Silver | 269 |
| Amylum | Alkaline matter | Red Litmus | 95 |
| | Acid matter | Blue Litmus | 95 |
| Antimonium Nigrum | Silica | Insolubility in HCl | 353 |
| Antimonii Oxidum | Higher oxides of antimony. | Insolubility in sol. of acid tartrate of potassium | 181 |
| Antimonium Tartaratum. | General [Sn] | Quantitative analysis | 579 |
| Aqua Aurantii Floris | Metallic matter (Pb, Cu, | Sulphuretted Hydrogen | 258 |
| Aqua Destillata | Fixed salts | Evaporation and ignition | 101 |
| | Tin, Lead, or Copper | Sulphuretted Hydrogen | 258 |
| | Calcium salts | Oxalate of Ammonium | 115 |
| | Chlorides | Nitrate of Silver | 269 |
| | Sulphates | Chloride or Nitrate of Barium | 311 |
| | Carbonates | Lime-water | 314 |
| Argenti Nitras | Other nitrates, etc. | Quantitative analysis, etc. | 584 |
| Argenti Oxidum | Metallic silver | Effervescence with Nitric acid | 216 |
| | General | Quantitative analysis | 584 |
| Argentum Purificatum. | Copper | Ammonia to the Nitric solution | 193 |
| Assafœtida | Earthy matter | Incineration | 101 |
| Atropia | Mineral matter | Incineration | 101 |
| Atropiæ Sulphas | Mineral matter | Incineration | 101 |
| Balsamum Peruvianum | Fixed oil | Immiscibility with Alcohol | 473 |
| | Alcohol | Non-diminution of volume when mixed with water | 425 |
| Bebeeriæ Sulphas | Mineral matter | Incineration | 101 |
| Bismuthi Carbonas | Nitrates (of Bismuth or Ammonium). | Sulphate of indigo | 290 |
| | Lead (carbonate) | Diluted Sulphuric acid | 214 |
| | Chloride (oxychloride of bismuth). | Nitrate of Silver | 269 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Bismuthi Oxidum | Moisture | Heated it loses weight | 596 |
| | Carbonate (of bismuth) | | |
| | Lead compounds | Sulphuric acid | 214 |
| | Chlorides | Nitrate of Silver | 269 |
| | Silver | HCl after elim. of BiCy AmHO | 218 |
| Bismuthi Subnitras | Lead (oxynitrate) | Diluted Sulphuric acid | 214 |
| | Chlorides (oxychloride of bismuth). | Nitrate of Silver | 269 |
| Bismuthum Purificatum. | Copper | Ammonia to Nitric solution | 193 |
| Bromum | General | Specific gravity (531) and boiling point | 515 |
| | Iodine | Starch | 275 |
| Cadmii Iodidum | Zinc (iodide) | Potash in excess, then Sulphydrate of Ammonium | 250 |
| | General | Quantitative analysis | 586 |
| Calcii Carbonas Præcipitata | Alumina, Oxide of Iron, and Phosphates. | Sacc. solution of lime to sol. in nitric acid | 107 |
| | Chlorides | Nitrate of Silver | 269 |
| Calcii Chloridum | Hypochlorite of Calcium | Hydrochloric acid | 292 |
| | Carbonic acid | Lime-water | 314 |
| Calcii Phosphas | Carbonates (of Calcium) | Effervescence with acids | 314 |
| | Alumina | Solution of potash | 139 |
| | Sand | Insolubility in acid | 353 |
| Calx | Carbonate of Calcium | Effervescence with acids | 314 |
| | Alumina, Oxide of Iron, etc. | Sacc. sol. of lime to sol. in acids | 107 |
| Calx Chlorata | General | Quantitative analysis | 565 |
| Cambogia | Rice, flour, sand, powdered bark. | Microscope, after exhausting with spt. and with cold water. | 403 |
| Camphora | Fixed salts | Non-volatility | 101 |
| Carbo Animalis Purificatus. | Earthy salts | Incineration (by help of red ox of mercury) | 101 |
| Carbo Ligni | More than 2 per ct. of ash | Incineration | 101 |
| Catechu Pallidum | Starch | Iodine | 275 |
| Cera Alba | Soft Fats | Melting-point | 517 |
| Cera Flava | Soft Fats | Melting-point | 517 |
| | Resin | Solubility in Alcohol | 464 |
| | Flour | Insolubility in Turpentine, Iodine | 275 |
| Cerii Oxalas | Carbonates and other oxalates. | Ash sol. in acids with effervescence | 314 |
| | Alumina | Insolubility of Hydrate in Ammonia | 230 |
| | General | More or less than 48 per cent. of ash | 229 |
| Cetaceum | Soft Fats | Melting-point | 517 |
| | Free acid | Litmus-paper | 95 |
| Chloral hydras | Other organic bodies | Sulphuric acid | 440 |
| | General | Quantitative analysis | 440 |
| | General | Specific gravity | 440 |
| Chloroform | General | Specific gravity | 532 |
| | Hydrocarbons | Sulphuric acid | 438 |
| | Non-volatile matter | Residue on evaporation | 101 |
| Copaiba | Wood-oil | Gelatinization at 270° F. | 471 |
| | | Incomplete solubility in Benzol | 471 |
| | | Oxidation | 444 |
| Creasotum | Carbolic acid | Non-volatility at 212° F. | 444 |
| | | Dextro-rotation of polarized ray | 444 |
| | | Crystallization on cooling | 444 |
| Cupri Sulphas | Iron (ferrous sulphate) | Nitric acid and ammonia | 160 |
| Elaterium | Carbonates (chalk) | Effervescence with acids | 314 |
| | General | Quantitative analysis | 418 |
| Fel Bovinum Purificatum. | Mucus, crude bile | Incomplete solubility in spirit | 457 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Ferri Arsenias | Sulphate (of sodium) | Chloride or Nitrate of Barium | 311 |
| | General | Quantitative analysis | 562 |
| Ferri Carbonas Saccharata | Sulphate (of ammonium) | Chloride or Nitrate of Barium | 311 |
| | General | Quantitative analysis | 562 |
| Ferri et Ammonii Citras | Tartrate (of iron and ammonium). | Ebullition with potash and saturation with Acetic acid ($=KHC_4H_4O_6$) | 322 |
| | General | Quantitative analysis | 578 |
| | Potassium or Sodium, salts | Alkalinity of ash | 101 |
| Ferri et Quiniæ Citras | Potassium or Sodium, salts | Alkalinity of ash | 101 |
| | General | Quantitative analysis | 578 |
| | Alkaloids other than Quinia. | Insolubility of ppt. alkaloid in ether | 603 |
| Ferri Oxidum Magneticum | Metallic iron | Effervescence with acids | 157 |
| | General | Quantitative analysis | 562 |
| Ferri Peroxidum Humidum | Ferrous hydrate | Red prussiate to acid solution | 160 |
| | Ferric oxhydrate | Insolubility in cold dil. Hydrochloric acid | 150 |
| Ferri Phosphas | Arsenicum (ferri arsenias). | Slip of Copper in acid solution | 172 |
| | General | Quantitative analysis | 562 |
| Ferri Sulphas | Ferric oxysulphate | Insolubility in Water | 143 |
| Ferri Sulphas Granulata | Ferric compounds | Ppt. of S in aqueous sol. by $H_2S$ | 160 |
| | Copper, etc. | Sulphuretted Hydrogen | 238 |
| Ferrum Redactum | Less than 50 per cent. | Quantitative analysis | 578 |
| Ferrum Tartaratum | Ferrous compounds | Red prussiate to acid solution | 160 |
| | Ammoniacal salts | Soda | 98 |
| | General | Quantitative analysis | 578 |
| Glycerina | General | Specific gravity | 448 |
| Hydrargyri Iodidum Rubrum. | Fixed Salts | Non-volatility | 208 |
| Hydrargyri Iodidum Viride. | Red Iodide | Insolubility in ether | 197 |
| Hydrargyri Oxidum Flavum. | Fixed salts (as NaCl) | Non-volatility | 208 |
| Hydrargyri Oxidum Rubrum | Fixed salts | Non-volatility | 208 |
| | Nitrates (of mercury) | Orange vapor on heating in a tube | 203 |
| Hydrargyri Subchloridum | Corrosive Sublimate | Treatment with ether | 202 |
| | Fixed salts | Non-volatility | 208 |
| Hydrargyri Sulphas | Fixed salts | Non-volatility | 208 |
| Hydrargyrum | Fixed metals (Pb, Sn, Zn, Bi, Cu). | Non-volatility | 208 |
| Hydrargyrum Ammoniatum. | Fixed salts | Non-volatility | 208 |
| Hydrargyrum cum Creta. | Mercuric oxide | Stannous chloride to sol. in HCl | 208 |
| Iodum | Fixed salts | Non-volatility | 274 |
| | Cyanide of Iodine | Physical characters | 274 |
| | General | Quantitative analysis | 564 |
| Jalapæ Resina | Resin | Solubility in Turpentine | 419 |
| Limonis Succus | Deficiency of Citric acid | Quantitative analysis | 325 |
| Liquor Ammoniæ | General | Specific gravity | 531 |
| | | Quantitative analysis | 547 |
| Liquor Ammoniæ Fortior | General impurity or deficiency. | Specific gravity and Quantitative analysis | 547 |
| | Carbonate of Ammonium | Lime-water | 314 |
| | Calcium salts (chloride, etc.). | Oxalate of Ammonium | 115 |
| | Iron salts (ferrous hydrate). | Sulphydrate of Ammonium | 160 |
| | Sulphur salts (AmHS) | Ammonio-sulphate of Copper | 303 |
| | Chloride of Ammonium | Nitrate of Silver to acidified sol. | 269 |
| | Sulphate of Ammonium | Chloride of Barium to acidified sol. | 311 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Liq. Antimonii Chloridi | . . . . . | . . . . . | 580 |
| Liq. Arsenicalis | . . . . . | . . . . . | 579 |
| Liq. Arsenici Hydrochlorici | General impurity or deficiency . . . . | Specific gravity . . . | 560 |
| Liq. Bismuthi et Ammonii Cit. | . . . . . | Quantitative analysis . | 560 |
| Liq. Calcis | . . . . . | . . . . . | 580 |
| Liq. Calcis Chloratæ | Deficiency in strength . | Quantitative analysis . . | 546 |
| Liq. Calcis Saccharatus | General impurity or deficiency . . . . | Specific gravity . . . . | 532 |
| | | Quantitative analysis . . | 546 |
| Liquor Chlori | General quality . . . . | Specific gravity . . . . | 532 |
| | Fixed matter . . . . | Residue on evaporation . | 101 |
| | Deficiency in strength . | Quantitative analysis . . | 564 |
| Liq. Ferri Perchloridi Fort. | Ferrous salts . . . . | Red Prussiate . . . . | 160 |
| Liq. Ferri Pernitratis | General impurity or deficiency . . . . | Specific gravity . . . . | 532 |
| | | Quantitative analysis . . | 578 |
| Liq. Ferri Persulphatis | . . . . . | . . . . . | |
| Liq. Hydrargyri Nitratis Acid. | Deficiency in strength . | Specific gravity . . . . | 532 |
| | Mercurous salts (nitrate) | Hydrochloric acid . . . | 207 |
| Liq. Lithii Effervescens. | General impurity or deficiency. | Quantitative analysis, etc. | 227 |
| Liq. Magnesii Carbonatis | Other magnesian salts . | Bitter taste ($MgCl_2MgSO_4$) | 119 |
| | General impurity or deficiency. | Quantitative analysis . . | 575 |
| Liq. Plumbi Subacetatis | General impurity or deficiency . . . . | Specific gravity . . . . | 532 |
| | | Quantitative analysis . . | 546 |
| Liquor Potassæ | General impurity or deficiency . . . . | Specific gravity . . . . | 532 |
| | | Quantitative analysis . . | 546 |
| | Carbonate (of potassium) . . . . . | Effervescence with acids . | 314 |
| | | Lime-water . . . . . | 314 |
| | Calcium salts . . . . | Oxalate of Ammonium . | 115 |
| | More than traces of: Silica . | Insol. in acid aft. evap. etc. | 353 |
| | More than traces of: Sulphates . | Chloride of Barium to acid sol. . . . . | 311 |
| | More than traces of: Chlorides . | Nitrate of silver to acid sol. . . . . | 269 |
| | More than traces of: Alumina . | Ammonia to acid sol. . . | 139 |
| Liq. Potassæ Effervescens | Deficiency in strength . | Quantitative analysis . . | 547 |
| | Bicarbonate of sodium . | Tartaric acid, etc. . . . | 571 |
| Liquor Sodæ | General impurity or deficiency . . . . | Specific gravity . . . | 532 |
| | | Quantitative analysis . . | 546 |
| | Calcium salts . . . . | Oxalate of Ammonium . | 115 |
| | Carbonate (of sodium) | Effervescence with acids . | 314 |
| | | Lime-water. . . . . | 314 |
| | More than traces of: Silica . | Insol. in acids after evap. | 353 |
| | More than traces of: Sulphates . | $BaCl_2$ to acid sol. . . . | 311 |
| | More than traces of: Chlorides . | $AgNO_3$ to acid sol. . . . | 269 |
| | More than traces of: Alumina . | Ammonia to acid sol. . . | 139 |
| Liquor Sodæ Chloratæ | Salts of Potassium or Ammonium. | Perchloride of Platinum to acid sol. . . . . | 98 |
| | General impurity or deficiency. | Quantitative analysis . . | 564 |
| Liq. Sodæ Effervescens | Calcium salts . . . . | Oxalate of Ammonium . | 115 |
| | Deficiency in strength . | Quantitative analysis . . | 546 |
| Lithii Carbonas | General impurity or deficiency. | Quantitative analysis . . [etc. | 227 |
| | Calcium salts . . . . | Oxalate of Ammonium, | 227 |
| | Alumina . . . . . | Lime-water, etc. . . . . | 227 |
| Lithii Citras | Deficiency in strength . | Quantitative analysis . . | 227 |
| Lycopodium | Starch, etc. . . . . | Microscope . . . . . | 403 |
| | More than 5 per cent. mineral matter. | Incineration . . . . . | 101 |
| | | Earthy matter sinks in $CS_2$ | |
| Magnesia | Carbonate of magnesium) | Effervescence with acids . | 119 |
| Magnesia Levis | Calcium (hydrate or carbonate). | Oxalate of Ammonium to acet. sol. . . . . | 115 |
| | Sulphates (of magnesium or sodium). | Chloride of Barium to acid sol. . . . . | 311 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Magnesii Carbonas | Alumina | Ammonia to acid sol. | 139 |
| | Sulphates (of magnesium or sodium). | Chloride of Barium to acid solution | 311 |
| Magnesii Carb. Levis | Calcium (carbonate) | Oxalic acid to Ammoniacal solution | 115 |
| | Iron, Lead, etc. | Sulph. hydrogen to sol. in acid + excess of Ammon. | 258 |
| | General impurity or deficiency. | Quantitative analysis | 575 |
| Magnesii Sulphas | Calcium (sulphate) | Oxalate of Ammonium | 115 |
| | Iron (sulphate) | Chlorinated Lime or Soda | 117 |
| | General impurities | Quantitative analysis | 575 |
| Manna | Deficiency of mannite | Quantitative analysis | 413 |
| Mel | Starch (flour) | Iodine | 275 |
| Morphiæ Hydrochloras | General impurities | Quantitative analysis | 607 |
| Olea Destillata | Fixed oil | Permanent greasy stain on paper | ... |
| | Alcohol | Loss in volume on shaking with water. | ... |
| Opium | Deficiency in morphia | Quantitative analysis | 607 |
| Plumbi Acetas | General | Quantitative analysis | 546 |
| Plumbi Carbonas | Sulphate of Lead or Barium or Silicates | Insolubility in Acetic acid (214, 103) | 353 |
| | Calcium (chalk) | Oxal. Ammonium after removing Lead | 115 |
| Plumbi Oxidum | Carbonates | Effervescence with acids | 314 |
| | Copper (oxide) | Ammonia to acid solution | 193 |
| Potassæ Causticæ | More than traces of Chloride | Nitrate of silver to acid solution | 269 |
| | More than traces of Sulphates | Chloride of Barium to acid solution | 311 |
| | General impurities (water, etc.). | Quantitative analysis | 546 |
| Potassæ Sulphuratæ | Excess of Carbonate or Sulphate | More than 25 per cent. insol. in spirit | 68 |
| Potassii Acetas | Iron and other metallic impurities | Sulphydrate of Ammonium | 258 |
| | Carbonate of Potassium | Effervescence with acids; alkalinity; insolubility in spirit | 314 68 |
| Potassii Bicarbonas | General | Quantitative analysis | 546 |
| Potassii Bromidum | Free bromine | Odor | 270 |
| | Iodide (of potassium) | Chlorine water and starch | 275 |
| | General | Quantitative analysis | 586 |
| Potassii Carbonas | More than traces of Silicate | Insol. in acid after evap. etc. | 353 |
| | More than traces of Sulphate | Chloride of Barium to acid solution | 311 |
| | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | General | Quantitative analysis | 546 |
| Potassii Chloras | Chloride (of potassium) | Nitrate of Silver | 269 |
| | Calcium (chloride) | Oxalate of Ammonium | 115 |
| Potassii Citras | General | Quantitative analysis | 547 |
| Potassii Ferridcyanidum. | Ferrocyanide (of potassium). | Ferric salt | 340 |
| Potassii Iodidum | Iodate (of potassium) | Tartaric acid and starch | 75 |
| | Chloride (of potassium) | Nitrate of Silver, etc. | 269 |
| | Carbonate (of potassium) | Sacc. solution of Lime. | 107 |
| Potassii Nitras | Sulphate (of potassium) | Chloride of Barium | 311 |
| | Chloride (of potassium) | Nitrate of Silver | 269 |
| Potassii Permanganas | General | Quantitative analysis | 558 |
| Potassii Sulphas | Acid Sulph. of Potassium | Test-paper | 95 |
| | Calcium (sulphate) | Oxalate of Ammonium | 115 |
| Potassii Sulphis | Hyposulphite (of potassium). | Sulphuric acid | 345 |
| Potassii Tartras Potassii Tart Acida | General | Quantitative analysis | 546 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Quiniæ Sulphas | Salicin | Sulphuric acid | 420 |
| | General | Quantitative analysis | 60 |
| Rhei Radix | Turmeric | Boracic acid | 334 |
| Santoninum | Mineral matter | Incineration | 101 |
| | Earthy soaps, etc. | Insolubility in spirit | 457 |
| Sapo Durus | Oil | Oily stain to paper | 457 |
| | Potassium compounds | Deliquescence of ash | 457 |
| Sapo Mollis | Earthy soaps, etc. | Insolubility in spirit | 457 |
| | Oil | Oily stain to paper | 457 |
| Scammoniæ Resina | Resin of guaiacum | Inner surface of Potato-paring | 419 |
| | Resin of Jalap | Insolubility in ether | 419 |
| Scammonium | Carbonate (of calcium and magnesium). | Effervescence with acids | 314 |
| | Starch (flour) | Solution of Iodine | 275 |
| | General impurities | Quantitative analysis | 421 |
| Sinapis | Starch (flour) | Solution of Iodine | 275 |
| Soda Caustica | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphate | Chloride of Barium to acid sol. | 311 |
| | General impurities (water, etc.). | Quantitative analysis | 546 |
| Soda Tartarata | General | Quantitative analysis | 546 |
| Soda Acetas | Acid or alkaline matter | Test-paper | 95 |
| | Sulphates (sodium or calcium). | Chloride of Barium to acid sol. | 311 |
| | Chlorides (sodium or calcium). | Nitrate of silver to acid sol. | 269 |
| Sodii Arsenias | Excess or deficiency of water of crystallization | Quantitative analysis | 536 |
| | General | Quantitative analysis | 556 |
| Sodii Bicarbonas | Neutral Carbonate (of sodium). | Mercuric chloride | 203 |
| | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphate | Chloride of Barium to acid sol. | 311 |
| | General | Quantitative analysis | 546 |
| Sodii Boras | General | Quantitative analysis | 546 |
| Sodii Carbonas | Excess or deficiency of water of cryst., etc. | Quantitative analysis | 596 |
| | More than traces of Chlorides | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphates | Chloride of Barium to acid sol. | 311 |
| Sodii Hyposulphis | General | Quantitative analysis | 560 |
| Sodii Nitras | Chloride (of sodium) | Nitrate of Silver | 269 |
| | Sulphate (of sodium) | Chloride or Nitrate of Barium | 311 |
| Sodii Phosphas | More than trace of Sulphate. | Chloride or Nitrate of Barium to acid sol. | 311 |
| | Deficiency or excess of water of cryst. | Quantitative analysis | 596 |
| Sodii Sulphas | Ammonium salts | Solut'n of potash heated | 98 |
| | Ferric salts | | 100 |
| | General | Quantitative analysis | 590 |
| | Excess or deficiency of water of cryst. | Quantitative analysis | 596 |
| Sodii Valerianas | Free soda or carbonate | Test-papers | 95 |
| | | Insolubility in spirit | 360 |
| Spiritus Ætheris Nitrosi | General | Specific gravity | 532 |
| | More than trace of acid | Effervescence with bicarbonate of sodium | 431 |
| | Free acid | More than "feeble" effervescence with bicarbonate of sodium | 314 |
| | Deficiency of nitrite of ethyl. | Quantitative analysis | 431 |
| Spiritus Ammoniæ Aromat. | General | Specific gravity | 532 |
| Spirit. Chloroformi | | | |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Magnesii Carbonas | Alumina | Ammonia to acid sol. | 139 |
| | Sulphates (of magnesium or sodium). | Chloride of Barium to acid solution | 311 |
| Magnesii Carb. Levis | Calcium (carbonate) | Oxalic acid to Ammoniacal solution | 115 |
| | Iron, Lead, etc. | Sulph. hydrogen to sol. in acid + excess of Ammon. | 258 |
| | General impurity or deficiency. | Quantitative analysis | 575 |
| Magnesii Sulphas | Calcium (sulphate) | Oxalate of Ammonium | 115 |
| | Iron (sulphate) | Chlorinated Lime or Soda | 117 |
| | General impurities | Quantitative analysis | 575 |
| Manna | Deficiency of mannite | Quantitative analysis | 413 |
| Mel | Starch (flour) | Iodine | 275 |
| Morphiæ Hydrochloras | General impurities | Quantitative analysis | 607 |
| Olea Destillata | Fixed oil | Permanent greasy stain on paper | ... |
| | Alcohol | Loss in volume on shaking with water. | ... |
| Opium | Deficiency in morphia | Quantitative analysis | 607 |
| Plumbi Acetas | General | Quantitative analysis | 546 |
| Plumbi Carbonas | Sulphate of Lead or Barium or Silicates | Insolubility in Acetic acid (214, 103) | 353 |
| | Calcium (chalk) | Oxal. Ammonium after removing Lead | 115 |
| Plumbi Oxidum | Carbonates | Effervescence with acids | 314 |
| | Copper (oxide) | Ammonia to acid solution | 193 |
| Potassæ Caustica | More than traces of Chloride | Nitrate of silver to acid solution | 269 |
| | More than traces of Sulphates | Chloride of Barium to acid solution | 311 |
| | General impurities (water, etc.). | Quantitative analysis | 546 |
| Potassæ Sulphuratæ | Excess of Carbonate or Sulphate | More than 25 per cent. insol. in spirit | 68 |
| Potassii Acetas | Iron and other metallic impurities | Sulphydrate of Ammonium | 258 |
| | Carbonate of Potassium | Effervescence with acids; alkalinity; insolubility in spirit | 314 68 |
| Potassii Bicarbonas | General | Quantitative analysis | 546 |
| Potassii Bromidum | Free bromine | Odor | 270 |
| | Iodide (of potassium) | Chlorine water and starch | 275 |
| | General | Quantitative analysis | 586 |
| Potassii Carbonas | More than traces of Silicate | Insol. in acid after evap. etc. | 353 |
| | More than traces of Sulphate | Chloride of Barium to acid solution | 311 |
| | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | General | Quantitative analysis | 546 |
| Potassii Chloras | Chloride (of potassium) | Nitrate of Silver | 269 |
| | Calcium (chloride) | Oxalate of Ammonium | 115 |
| Potassii Citras | General | Quantitative analysis | 547 |
| Potassii Ferridcyanidum. | Ferrocyanide (of potassium). | Ferric salt | 340 |
| Potassii Iodidum | Iodate (of potassium) | Tartaric acid and starch | 75 |
| | Chloride (of potassium) | Nitrate of Silver, etc. | 269 |
| | Carbonate (of potassium) | Sacc. solution of Lime. | 107 |
| Potassii Nitras | Sulphate (of potassium) | Chloride of Barium | 311 |
| | Chloride (of potassium) | Nitrate of Silver | 269 |
| Potassii Permanganas | General | Quantitative analysis | 558 |
| Potassii Sulphas | Acid Sulph. of Potassium | Test-paper | 95 |
| | Calcium (sulphate) | Oxalate of Ammonium | 115 |
| Potassii Sulphis | Hyposulphite (of potassium). | Sulphuric acid | 345 |
| Potassii Tartras Potassii Tart Acida | General | Quantitative analysis | 546 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Quinia Sulphas | Salicin | Sulphuric acid | 420 |
| | General | Quantitative analysis | 60 |
| Rhei Radix | Turmeric | Boracic acid | 334 |
| Santoninum | Mineral matter | Incineration | 101 |
| | Earthy soaps, etc. | Insolubility in spirit | 457 |
| Sapo Durus | Oil | Oily stain to paper | 457 |
| | Potassium compounds | Deliquescence of ash | 457 |
| Sapo Mollis | Earthy soaps, etc. | Insolubility in spirit | 457 |
| | Oil | Oily stain to paper | 457 |
| Scammonia Resina | Resin of guaiacum | Inner surface of Potato-pariug | 419 |
| | Resin of Jalap | Insolubility in ether | 419 |
| Scammonium | Carbonate (of calcium and magnesium). | Effervescence with acids | 314 |
| | Starch (flour) | Solution of Iodine | 275 |
| | General impurities | Quantitative analysis | 421 |
| Sinapis | Starch (flour) | Solution of Iodine | 275 |
| Soda Caustica | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphate | Chloride of Barium to acid sol. | 311 |
| | General impurities (water, etc.). | Quantitative analysis | 546 |
| Soda Tartarata | General | Quantitative analysis | 546 |
| Soda Acetas | Acid or alkaline matter | Test-paper | 93 |
| | Sulphates (sodium or calcium). | Chloride of Barium to acid sol. | 311 |
| | Chlorides (sodium or calcium). | Nitrate of silver to acid sol. | 269 |
| Sodii Arsenias | Excess or deficiency of water of crystallization | Quantitative analysis | 556 |
| | General | Quantitative analysis | 556 |
| Sodii Bicarbonas | Neutral Carbonate (of sodium). | Mercuric chloride | 203 |
| | More than traces of Chloride | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphate | Chloride of Barium to acid sol. | 311 |
| | General | Quantitative analysis | 546 |
| Sodii Boras | General | Quantitative analysis | 546 |
| Sodii Carbonas | Excess or deficiency of water of cryst., etc. | Quantitative analysis | 596 |
| | More than traces of Chlorides | Nitr. of Silver to acid sol. | 269 |
| | More than traces of Sulphates | Chloride of Barium to acid sol. | 311 |
| Sodii Hyposulphis | General | Quantitative analysis | 560 |
| Sodii Nitras | Chloride (of sodium) | Nitrate of Silver | 269 |
| | Sulphate (of sodium) | Chloride or Nitrate of Barium | 311 |
| Sodii Phosphas | More than trace of Sulphate. | Chloride or Nitrate of Barium to acid sol. | 311 |
| | Deficiency or excess of water of cryst. | Quantitative analysis | 596 |
| Sodii Sulphas | Ammonium salts | Solut'n of potash heated | 93 |
| | Ferric salts | | 100 |
| | General | Quantitative analysis | 590 |
| | Excess or deficiency of water of cryst. | Quantitative analysis | 596 |
| Sodii Valerianas | Free soda or carbonate | Test-papers | 95 |
| | | Insolubility in spirit | 360 |
| Spiritus Ætheris Nitrosi | General | Specific gravity | 532 |
| | More than trace of acid | Effervescence with bicarbonate of sodium | 431 |
| | Free acid | More than "feeble" effervescence with bicarbonate of sodium | 314 |
| | Deficiency of nitrite of ethyl. | Quantitative analysis | 431 |
| Spiritus Ammoniæ Aromat. Spirit. Chloroformi | General | Specific gravity | 532 |

| Name of Preparation. | Impurities. | Tests. | Page. |
|---|---|---|---|
| Spiritus Rectificatus | General (excess of water) | Specific gravity | 531 |
| | Resin or oil | Opalescence on dilution | 425 |
| | More than traces of fusel oil, etc. | Nitrate of Silver | ... |
| Spiritus Tenuior | General (excess of water) | Specific gravity | 423 |
| Strychnia | Brucia | Nitric acid | 391 |
| | Mineral matter | Incineration | 101 |
| Sulphur Præcipitatum | Sulphate of calcium | Appearance under microscope | 304 |
| | | Residue on ignition | ... |
| Sulphur Sublimatum | Earthy matter | Incineration | 101 |
| | Trace of Acid ($H_2SO_4$ or $H_2SO_3$). | Litmus-paper | 95 |
| | Sulphide of Arsenicum | Ammonia | 176 |
| Sulphuris Iodidum | Deficiency of Iodine | Quantitative analysis | 275 |
| Syrupi | Deficiency of Sugar | Specific gravity | 532 |
| Tamarindus | Traces of Copper | Iron | 192 |
| Veratria | Mineral matter | Incineration | 101 |
| Zinci Acetas | Sulphates | Chloride or Nitrate of Barium | 311 |
| | Chlorides | Nitrate of Silver | 269 |
| | Metals (As, Cd, Cu, Pb) | Sulphuretted Hydrogen | 238 |
| | Iron (acetate) | Nitric Acid: Ammonia | 162 |
| | Copper (acetate) | Ammonia | 193 |
| Zinci Carbonas | Sulphates | Chloride or Nitrate of Barium to acid sol. | 311 |
| | Chlorides | Nitr. of Silver to acid sol. | 269 |
| | Copper (carbonate) | Ammonia to acid solution | 193 |
| | Metals (As, Cd, Cu, Pb) | Sulphuretted Hydrogen | 238 |
| Zinci Chloridum | Sulphates | Chloride or Nitrate of Barium | 311 |
| | Calcium (chloride) | Oxalate of Ammonium | 115 |
| | Ferrous salts (chloride) | Ferridcyanide of Potass'm | 160 |
| | Ferric salts (chloride) | Ferrocyanide of Potass'm | 160 |
| Zinci Oxidum | Carbonate (of zinc) | Effervescence with acids | 314 |
| | Sulphates (sodium or zinc). | Chloride or Nitrate of Barium to acid sol. | 311 |
| | Chlorides | Nitr. of Silver to acid sol. | 269 |
| | Copper (oxide) | Ammonia to acid solution | 193 |
| Zinci Sulphas | Metal (As, Cd, Cu, Pb) | Sulphuretted Hydrogen | 238 |
| | Iron (sulphate) | Tincture of galls | 356 |
| | | Nitric acid: Ammonia | 160 |
| | Copper (sulphate) | Ammonia | 193 |
| Zinci Valerianas | Sulphate (of zinc) | Chloride or Nitrate of Barium | 311 |
| | Butyrate (of zinc) | Acetate of Copper, etc. | 361 |

**Equivalent weights** of Citric Acid, Tartaric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium, Bicarbonate of Sodium, Carbonate of Ammonium, and Carbonate of Magnesium; repeated (in **black**) for 20 parts of each, and incidentally (in roman) for other proportions. (Exact to two places of decimals.)

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| Citric Acid ($H_3C_6H_5O_7H_2O$) ÷ 3 × 2 = 140 . . . . . | **20.00** | 18.66 | **16.96** | **14.00** | **9.78** | **16.66** | **23.72** | **29.31** |
| Tartaric Acid $H_2C_4H_4O_6$ = 150 . . . . . | 21.43 | **20.00** | **18.26** | **15.00** | **10.49** | **17.85** | **25.42** | **31.41** |
| Carbonate of Potassium $K_2CO_3 + 16$ % Aq. = 164.285 . | **23.47** | **21.90** | **20.00** | 16.43 | 11.48 | 19.52 | 27.87 | 34.40 |
| Bicarbonate of Potassium $2(KHCO_3)$ = 200 . . . . . | **28.57** | **26.66** | 24.34 | **20.00** | 13.98 | 23.81 | 33.89 | 41.90 |
| Carbonate of Sodium $Na_2CO_3,10H_2O$ = 286 . . . . . | **40.08** | **38.13** | 34.81 | 28.60 | **20.00** | 34.04 | 48.47 | 59.98 |
| Bicarbonate of Sodium $2(NaHCO_3)$ = 168 . . . . . | **24.00** | **22.40** | 20.45 | 16.80 | 11.74 | **20.00** | 28.47 | 35.18 |
| Carbonate of Ammonium ($N_4H_{16}C_2O_6$) ÷ 2 = 118 . . . | **16.85** | **15.73** | 14.36 | 11.80 | 8.25 | 14.04 | **20.00** | 24.71 |
| Carb. of Magnes. ($(MgCO_3)_3,Mg2HO,4H_2O$) ÷ 4 = 95.5 | **13.64** | **12.73** | 11.62 | 9.55 | 6.68 | 11.37 | 16.18 | **20.00** |

The amount of acid given in any column will saturate the amount of carbonate in the same column, and *vice versâ*.
The amounts of carbonate in any column are equal to each other in chemical power.
Lemon-juice (sp. gr. 1.039) contains, on an average, 7 per cent. by weight of citric acid.

*The same Table in round numbers, for purposes of prescribing and dispensing.*

(The old names in Latin.)

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| Citric Acid . . . . . | **20** | 19 | **17** | **14** | **10** | **17** | **24** | **30** |
| Tartaric acid . . . . . | 22 | **20** | **18** | **15** | **11** | **18** | **26** | **32** |
| Carbonate of Potassium (Potassæ Carbonas) . . . . . | **24** | **22** | **20** | 16 | 12 | 20 | 28 | 35 |
| Bicarbonate of Potassium (Potassæ Bicarbonas) . . . . | **29** | **27** | 24 | **20** | 14 | 24 | 34 | 42 |
| Barbonate of Sodium (cryst.) (Sodæ Carbonas) . . . . | **40** | **38** | 35 | 28 | **20** | 34 | 49 | 60 |
| Bicarbonate of Sodium (Sodæ Bicarbonas) . . . . . | **24** | **22** | 20 | 17 | 12 | **20** | 29 | 36 |
| Carbonate of Ammonium (Ammoniæ Carbonas) . . . | **17** | **16** | 14 | 12 | 8 | 14 | **20** | 25 |
| Carbonate of Magnesium (Magnesiæ Carbonas) . . . . | **13** | **13** | 11 | 9 | 7 | 11 | 16 | **20** |

The Table is read thus: 20 grains of Citric Acid will saturate 29 grains of Bicarbonate of Potassium; 20 grains of Bicarbonate of Sodium will saturate, or be saturated by, 18 grains of Tartaric Acid; 11 grains of Tartaric Acid = 8 grains of Carbonate of Ammonium; 20 grains of Bicarbonate of Sodium are equivalent to, or will do as much work as, 34 grains of Carbonate of Sodium; 14 grains of Citric Acid are as strong as 15 of Tartaric Acid. It is occasionally convenient to double the numbers, halve them, or take some other proportion; also to employ them in weights other than grains.

Lemon-juice contains, on an average, 32½ grains of citric acid in 1 fluidounce, or 4 grains per fluidrachm.

TABLE OF THE PERCENTAGE OF SULPHURIC ACID ($H_2SO_4$), THE SULPHURIC RADICAL ($SO_4$), AND SULPHURIC ANHYDRIDE ($SO_3$), IN DILUTE SULPHURIC ACID OF DIFFERENT SPECIFIC GRAVITIES.*

| Specific gravity. | $H_2SO_4$. | $SO_4$. | $SO_3$. |
|---|---|---|---|
| 1.8485 | 100 | 97.848 | 81.54 |
| 1.8475 | 99 | 96.864 | 80.72 |
| 1.8460 | 98 | 95.880 | 79.90 |
| 1.8439 | 97 | 94.908 | 79.09 |
| 1.8430 | 96.8 | 94.82 | 79.02 |
| 1.8410 | 95 | 93.936 | 78.28 |
| 1.8376 | 95 | 92.952 | 77.46 |
| 1.8336 | 94 | 91.980 | 76.65 |
| 1.8290 | 93 | 90.996 | 75.83 |
| 1.8233 | 92 | 90.024 | 75.02 |
| 1.8179 | 91 | 89.040 | 74.20 |
| 1.8115 | 90 | 88.068 | 73.39 |
| 1.8043 | 89 | 87.084 | 72.57 |
| 1.7962 | 88 | 86.100 | 71.75 |
| 1.7870 | 87 | 85.128 | 70.94 |
| 1.7774 | 86 | 84.144 | 70.12 |
| 1.7673 | 85 | 83.172 | 69.31 |
| 1.7570 | 84 | 82.188 | 68.49 |
| 1.7465 | 83 | 81.216 | 67.68 |
| 1.7360 | 82 | 80.232 | 66.86 |
| 1.7245 | 81 | 79.260 | 66.05 |
| 1.7120 | 80 | 78 276 | 65.23 |
| 1.6993 | 79 | 77.304 | 64.42 |
| 1.6870 | 78 | 76.320 | 63.60 |
| 1.6750 | 77 | 75.336 | 62.78 |
| 1.6630 | 76 | 74.364 | 61.97 |
| 1.6520 | 75 | 73.380 | 61.15 |
| 1.6415 | 74 | 72.408 | 60.34 |
| 1.6321 | 73 | 71.424 | 59.52 |
| 1.6204 | 72 | 70.452 | 58.71 |
| 1.6090 | 71 | 69.468 | 57.89 |
| 1.5975 | 70 | 68.496 | 57.08 |
| 1.5868 | 69 | 67.512 | 56 26 |
| 1.5760 | 68 | 66.540 | 55.45 |
| 1.5648 | 67 | 65.556 | 54.63 |
| 1.5503 | 66 | 64.584 | 53.82 |
| 1.5390 | 65 | 63.600 | 53.00 |
| 1.5280 | 64 | 62.616 | 52.18 |

* The practice of regarding anhydrides as acids, and aqueous solutions of the alkalies as containing anhydrous oxides, is being rapidly abandoned; the old Tables of the strength of acid and alkaline solutions possess, consequently, diminished value. The publication, therefore, of Tables founded on those of Dalton, Ure, Carius, and J. Watts, but showing the percentage of true acid and acidulous radical as well as anhydride, and of alkaline hydrate and metal as well as oxide, will, it is hoped, prove useful to chemists generally as well as students.

| Specific gravity. | $H_2SO_4$. | $SO_4$. | $SO_3$. |
|---|---|---|---|
| 1.5170 | 63 | 61.644 | 51.37 |
| 1.5066 | 62 | 60.660 | 50.55 |
| 1.4960 | 61 | 59.688 | 49.74 |
| 1.4860 | 60 | 58.704 | 48.92 |
| 1.4760 | 59 | 57.732 | 48.11 |
| 1.4660 | 58 | 56.748 | 47.29 |
| 1.4560 | 57 | 55.776 | 46.48 |
| 1.4460 | 56 | 54.792 | 45.66 |
| 1.4360 | 55 | 53.820 | 44.85 |
| 1.4265 | 54 | 52 836 | 44 03 |
| 1.4170 | 53 | 51.864 | 43.22 |
| 1.4073 | 52 | 50.880 | 42.40 |
| 1.3977 | 51 | 49.896 | 41.58 |
| 1.3884 | 50 | 48.924 | 40.77 |
| 1.3788 | 49 | 47.940 | 39.95 |
| 1.3697 | 48 | 46 968 | 39.14 |
| 1.3612 | 47 | 45.984 | 38.32 |
| 1.3530 | 46 | 45.012 | 37.51 |
| 1.3440 | 45 | 44.028 | 36.69 |
| 1.3345 | 44 | 43.056 | 35.88 |
| 1.3255 | 43 | 42.072 | 35.06 |
| 1.3165 | 42 | 41.100 | 34.25 |
| 1.3080 | 41 | 40.116 | 33.43 |
| 1.2999 | 40 | 39.132 | 32.61 |
| 1.2913 | 39 | 38.160 | 31.80 |
| 1.2826 | 38 | 37.176 | 30.98 |
| 1.2740 | 37 | 36 204 | 30.17 |
| 1.2654 | 36 | 35.220 | 29.35 |
| 1.2572 | 35 | 34 248 | 28.54 |
| 1.2490 | 34 | 33.264 | 27.72 |
| 1.2409 | 33 | 32.292 | 26.91 |
| 1.2334 | 32 | 31.308 | 26.09 |
| 1.2260 | 31 | 30.336 | 25.28 |
| 1.2184 | 30 | 29.352 | 24.46 |
| 1.2108 | 29 | 28.380 | 23.65 |
| 1.2032 | 28 | 27.396 | 22.83 |
| 1.1956 | 27 | 26.412 | 22.01 |
| 1.1876 | 26 | 25.440 | 21.20 |
| 1.1792 | 25 | 24.456 | 20.38 |
| 1.1706 | 24 | 23.484 | 19.57 |
| 1.1626 | 23 | 22.500 | 18.75 |
| 1.1549 | 22 | 21.528 | 17.94 |
| 1.1480 | 21 | 20.544 | 17.12 |
| 1.1410 | 20 | 19.572 | 16.31 |
| 1.1330 | 19 | 18.588 | 15.49 |
| 1.1246 | 18 | 17.616 | 14.68 |
| 1.1165 | 17 | 16.632 | 13.86 |
| 1.1090 | 16 | 15.660 | 13.05 |
| 1.1019 | 15 | 14.676 | 12.23 |
| 1.0953 | 14 | 13.692 | 11.41 |
| 1.0940 | 13.65 | 13.370 | 11.14 |
| 1.0887 | 13 | 12.720 | 10.60 |

| Specific gravity. | $H_2SO_4$. | $SO_4$. | $SO_3$. |
|---|---|---|---|
| 1.0809 | 12 | 11.736 | 9.78 |
| 1.0743 | 11 | 10.764 | 8.97 |
| 1.0682 | 10 | 9.780 | 8.15 |
| 1.0614 | 9 | 8.808 | 7.34 |
| 1.0544 | 8 | 7.824 | 6.52 |
| 1.0477 | 7 | 6.852 | 5.71 |
| 1.0405 | 6 | 5.868 | 4.89 |
| 1.0336 | 5 | 4.896 | 4.08 |
| 1.0268 | 4 | 3.912 | 3 26 |
| 1.0206 | 3 | 2.9352 | 2.446 |
| 1.0140 | 2 | 1.956 | 1.63 |
| 1.0074 | 1 | 0.9785 | 0.8154 |

$H_2SO_4 = 98$; $SO_4 = 96$; $SO_3 = 80$.

---

TABLE OF THE PERCENTAGE OF NITRIC ACID ($HNO_3$), THE NITRIC RADICAL ($NO_3$), AND NITRIC ANHYDRIDE ($N_2O_5$), IN DILUTE NITRIC ACID OF DIFFERENT SPECIFC GRAVITIES.

| Specific gravity. | $HNO_3$. | $NO_3$. | $N_2O_5$. |
|---|---|---|---|
| 1.5000 | 92.988 | 91.507 | 79.700 |
| 1.4980 | 92.053 | 90.592 | 78 903 |
| 1.4960 | 91.124 | 89.677 | 78.106 |
| 1.4940 | 90.194 | 88.762 | 77.309 |
| 1.4910 | 89.264 | 87 847 | 76.512 |
| 1.4880 | 88.334 | 86.932 | 75.715 |
| 1.4850 | 87.404 | 86.017 | 74.918 |
| 1.4820 | 86.484 | 85.102 | 74.121 |
| 1.4790 | 85.545 | 84 187 | 73.324 |
| 1.4760 | 84.615 | 83.272 | 72.527 |
| 1.4730 | 83.680 | 82.357 | 71.730 |
| 1.4700 | 82.755 | 81.442 | 70.933 |
| 1.4670 | 81.825 | 80.527 | 70.136 |
| 1.4640 | 80.895 | 79.611 | 69.339 |
| 1.4600 | 79.966 | 78.696 | 68.542 |
| 1.4570 | 79.036 | 77.781 | 67.745 |
| 1.4530 | 78.106 | 76.866 | 66.948 |
| 1.4500 | 77.181 | 75.956 | 66.155 |
| 1.4460 | 76.246 | 75.036 | 65.354 |
| 1.4424 | 75.316 | 74.121 | 64.557 |
| 1.4385 | 74.386 | 73.206 | 63.760 |
| 1.4346 | 73.457 | 72.291 | 62 963 |
| 1.4306 | 72.527 | 71.376 | 62.166 |
| 1.4269 | 71.587 | 70.461 | 61.369 |
| 1.4228 | 70.667 | 69.546 | 60.572 |
| 1.4200 | 70.0 | 68.889 | 60.0 |
| 1.4189 | 69.737 | 68.631 | 59.775 |
| 1.4147 | 68.807 | 67.715 | 58.978 |
| 1.4107 | 67.878 | 66.800 | 58.181 |

| Specific gravity. | $HNO_3$. | $NO_2$. | $N_2O_5$. |
|---|---|---|---|
| 1.4065 | 66.948 | 65 885 | 57.884 |
| 1.4023 | 66.018 | 64.970 | 56.587 |
| 1.3978 | 65.090 | 64.055 | 55.790 |
| 1.3945 | 64.156 | 63.140 | 54.993 |
| 1.3882 | 63 229 | 62 225 | 54.196 |
| 1.3833 | 62.299 | 61.310 | 53.399 |
| 1.3783 | 61.369 | 60.395 | 52.602 |
| 1.3732 | 60.439 | 59.480 | 51.805 |
| 1.3681 | 59.579 | 58.634 | 51.068 |
| 1.3630 | 58.579 | 57.650 | 50.211 |
| 1.3579 | 57.650 | 56.734 | 49.414 |
| 1.3529 | 56.720 | 55.819 | 48.617 |
| 1.3477 | 55.790 | 54.904 | 47.820 |
| 1.3427 | 54.860 | 53.989 | 47.023 |
| 1.3376 | 53.930 | 53.074 | 46.226 |
| 1.3323 | 53.000 | 52.159 | 45.429 |
| 1.3270 | 52.069 | 51.244 | 44.632 |
| 1.3216 | 51.141 | 50.329 | 43.835 |
| 1 3163 | 50.211 | 49.414 | 43.038 |
| 1.3110 | 49.281 | 48.499 | 42.241 |
| 1.3056 | 48.355 | 47.587 | 41.447 |
| 1.3001 | 47.180 | 46.439 | 40.440 |
| 1.2947 | 46.299 | 45.564 | 39.685 |
| 1.2887 | 45.562 | 44.838 | 39.053 |
| 1.2826 | 44.632 | 43.923 | 38.256 |
| 1 2765 | 43.702 | 43.008 | 37.459 |
| 1.2705 | 42.772 | 42.095 | 36.662 |
| 1.2644 | 41.842 | 41.178 | 35.865 |
| 1.2583 | 40.913 | 40.263 | 35.068 |
| 1.2523 | 39.983 | 39.348 | 34.271 |
| 1.2462 | 39.053 | 38.433 | 33.474 |
| 1.2402 | 38.123 | 37.888 | 32.677 |
| 1.2341 | 37.190 | 36.603 | 31.880 |
| 1 2277 | 36.263 | 35.694 | 31.083 |
| 1.2212 | 35.334 | 34.735 | 30.286 |
| 1.2148 | 34.404 | 33.857 | 29.489 |
| 1.2084 | 33.474 | 32.942 | 28.692 |
| 1.2019 | 32.544 | 32.027 | 27.895 |
| 1.1958 | 31.614 | 31.112 | 27.098 |
| 1.1895 | 30.684 | 30.197 | 26.301 |
| 1.1833 | 29 755 | 29.282 | 25.504 |
| 1.1770 | 28.825 | 28.367 | 24.707 |
| 1.1709 | 27.890 | 27.454 | 23.910 |
| 1.1648 | 26.965 | 26.537 | 23.113 |
| 1.1587 | 26.035 | 25.622 | 22.316 |
| 1.1526 | 25.105 | 24.707 | 21.519 |
| 1.1465 | 24.176 | 23.792 | 20.722 |
| 1.1403 | 23.246 | 23.877 | 19.925 |
| 1.1345 | 22.316 | 21.962 | 19.128 |
| 1.1286 | 21.386 | 21.047 | 18.331 |
| 1.1227 | 20.456 | 20.131 | 17.534 |
| 1.1168 | 19.526 | 19.217 | 16.737 |

| Specific gravity. | $HNO_3$. | $NO_3$. | $N_2O_5$. |
|---|---|---|---|
| 1.1109 | 18.597 | 18.301 | 15.940 |
| 1.1051 | 17 667 | 17.386 | 15.143 |
| 1.1010 | 17.442 | 17.165 | 14.950 |
| 1.0993 | 16.737 | 16.472 | 14.346 |
| 1.0935 | 15.807 | 15.562 | 13.549 |
| 1.0878 | 14.877 | 14.634 | 12.752 |
| 1.0821 | 13.947 | 13.726 | 11.955 |
| 1.0764 | 13.017 | 12.811 | 11.158 |
| 1.0708 | 12.088 | 11.922 | 10.361 |
| 1.0651 | 11.158 | 10.981 | 9.564 |
| 1.0595 | 10.228 | 10.066 | 8.767 |
| 1.0540 | 9.298 | 9.151 | 7.970 |
| 1.0485 | 8.368 | 8.235 | 7.173 |
| 1.0430 | 7.439 | 7.320 | 6.376 |
| 1.0375 | 6.508 | 6.405 | 5.579 |
| 1.0320 | 5.579 | 5.490 | 4.782 |
| 1.0267 | 4.649 | 4.575 | 3.985 |
| 1.0212 | 3 719 | 3.660 | 3.188 |
| 1.0159 | 2.789 | 2.745 | 2.391 |
| 1.0106 | 1.860 | 1.830 | 1.594 |
| 1.0053 | 0.930 | 0.915 | 0.797 |

$HNO_3 = 63$; $NO_3 = 62$; $N_2O_5 \div 2 = 54$.

---

TABLE OF THE QUANTITY OF HYDROCHLORIC ACID IN THE LIQUID ACID OF DIFFERENT SPECIFIC GRAVITIES (URE).

| Specific gravity. | Acid of 1.2 in 100. | Hydrochloric acid gas. | Chlorine. |
|---|---|---|---|
| 1.2000 | 100 | 40.777 | 39.675 |
| 1.1982 | 99 | 40.369 | 39.278 |
| 1.1964 | 98 | 39.961 | 38.882 |
| 1.1946 | 97 | 39.554 | 38.485 |
| 1.1928 | 96 | 39.146 | 38.089 |
| 1.1910 | 95 | 38.738 | 37.692 |
| 1.1893 | 94 | 38.330 | 87 296 |
| 1.1875 | 93 | 37.923 | 36.900 |
| 1.1857 | 92 | 37.516 | 36.503 |
| 1.1846 | 91 | 37.108 | 36.107 |
| 1.1822 | 90 | 36.700 | 35.707 |
| 1.1802 | 89 | 36.292 | 35.310 |
| 1.1782 | 88 | 35.884 | 34.913 |
| 1.1762 | 87 | 35.476 | 34.517 |
| 1.1741 | 86 | 35.068 | 34.121 |
| 1.1721 | 85 | 34.660 | 33.724 |
| 1.1701 | 84 | 34.252 | 33.328 |
| 1.1681 | 83 | 33.845 | 32 931 |
| 1.1661 | 82 | 33.437 | 32.535 |
| 1.1641 | 81 | 33.029 | 32.136 |
| 1.1620 | 80 | 32.621 | 31.743 |

| Specific gravity. | Acid of 1.2 in 100. | Hydrochloric acid gas. | Chlorine. |
|---|---|---|---|
| 1.1599 | 79 | 32.213 | 31.348 |
| 1.1578 | 78 | 31.805 | 30.946 |
| 1.1557 | 77 | 31.398 | 30.550 |
| 1.1536 | 76 | 30.990 | 30.153 |
| 1.1515 | 75 | 30.582 | 29.757 |
| 1.1494 | 74 | 30.174 | 29.361 |
| 1.1473 | 73 | 29.767 | 28.964 |
| 1.1452 | 72 | 29.359 | 28.567 |
| 1.1431 | 71 | 28.951 | 28.171 |
| 1.1410 | 70 | 28.544 | 27.772 |
| 1.1389 | 69 | 28.136 | 27.376 |
| 1.1369 | 68 | 27.728 | 26.979 |
| 1.1349 | 67 | 27.321 | 26.583 |
| 1.1328 | 66 | 26.913 | 26.186 |
| 1.1308 | 65 | 26.508 | 25.789 |
| 1.1287 | 64 | 26.098 | 25.392 |
| 1.1267 | 63 | 25.690 | 24.996 |
| 1.1247 | 62 | 25.282 | 24.599 |
| 1.1226 | 61 | 24.874 | 24.202 |
| 1.1206 | 60 | 24.466 | 23.805 |
| 1.1185 | 59 | 24.058 | 23.408 |
| 1.1164 | 58 | 23.650 | 23.012 |
| 1.1143 | 57 | 23.242 | 22.615 |
| 1.1123 | 56 | 22 834 | 22.218 |
| 1.1102 | 55 | 22.426 | 21.822 |
| 1.1082 | 54 | 22.019 | 21.425 |
| 1.1061 | 53 | 21.611 | 21.028 |
| 1.1041 | 52 | 21.203 | 20.632 |
| 1.1020 | 51 | 20.796 | 20.235 |
| 1.1000 | 50 | 20.388 | 19.837 |
| 1.0980 | 49 | 19.980 | 19.440 |
| 1.0960 | 48 | 19.572 | 19.044 |
| 1.0939 | 47 | 19.165 | 18.647 |
| 1.0919 | 46 | 18.757 | 18.250 |
| 1.0899 | 45 | 18.349 | 17.854 |
| 1.0879 | 44 | 17.941 | 17.457 |
| 1.0859 | 43 | 17.584 | 17.060 |
| 1.0838 | 42 | 17.126 | 16.664 |
| 1.0818 | 41 | 16.718 | 16.267 |
| 1.0798 | 40 | 16.310 | 15.870 |
| 1.0778 | 39 | 15.902 | 15.474 |
| 1.0758 | 38 | 15.494 | 15.077 |
| 1.0738 | 37 | 15.087 | 14.680 |
| 1.0718 | 36 | 14.679 | 14.284 |
| 1.0697 | 35 | 14.271 | 13.887 |
| 1.0677 | 34 | 13.863 | 13.490 |
| 1.0657 | 33 | 13.456 | 13.094 |
| 1.0637 | 32 | 13.049 | 12.597 |
| 1.0617 | 31 | 12.641 | 12.300 |
| 1.0597 | 30 | 12.233 | 11.903 |
| 1.0577 | 29 | 11.825 | 11.506 |
| 1.0557 | 28 | 11.416 | 11.109 |

| Specific gravity. | Acid of 1.2 in 100. | Hydrochloric acid gas. | Chlorine. |
|---|---|---|---|
| 1.0537 | 27 | 11.010 | 10.712 |
| 1.0517 | 26 | 10.602 | 10.316 |
| 1.0497 | 25 | 10.194 | 9.919 |
| 1.0477 | 24 | 9.786 | 9.522 |
| 1.0457 | 23 | 9.379 | 9.126 |
| 1.0437 | 22 | 8.971 | 8.729 |
| 1.0417 | 21 | 8.563 | 8.332 |
| 1.0397 | 20 | 8.155 | 7 935 |
| 1.0377 | 19 | 7.747 | 7.538 |
| 1.0357 | 18 | 7.340 | 7.141 |
| 1.0337 | 17 | 6.932 | 6.745 |
| 1.0318 | 16 | 6.524 | 6.348 |
| 1.0298 | 15 | 6.116 | 5.951 |
| 1.0279 | 14 | 5.709 | 5.554 |
| 1.0259 | 13 | 5.301 | 5.158 |
| 1.0239 | 12 | 4.893 | 4.762 |
| 1.0220 | 11 | 4.486 | 4.365 |
| 1.0200 | 10 | 4.078 | 3.998 |
| 1.0180 | 9 | 3.670 | 3.571 |
| 1.0160 | 8 | 3.262 | 3.174 |
| 1.0140 | 7 | 2.854 | 2.778 |
| 1.0120 | 6 | 2.447 | 2.381 |
| 1.0110 | 5 | 2.039 | 1.984 |
| 1.0080 | 4 | 1.631 | 1.588 |
| 1.0060 | 3 | 1.224 | 1.191 |
| 1.0040 | 2 | 0.816 | 0.795 |
| 1.0020 | 1 | 0.408 | 0.397 |

TABLE OF THE PERCENTAGE OF PHOSPHORIC ACID ($H_3PO_4$), THE PHOSPHORIC RADICAL ($PO_4$), AND PHOSPHORIC ANHYDRIDE ($P_2O_5$), IN AQUEOUS SOLUTIONS OF PHOSPHORIC ACID OF DIFFERENT SPECIFIC GRAVITIES.

| Sp. gr. | $H_3PO_4$. | $PO_4$. | $P_2O_5$. |
|---|---|---|---|
| 1.508 | 68.45 | 66.37 | 49.60 |
| 1.492 | 66.82 | 64.77 | 48.41 |
| 1.476 | 65.01 | 63.02 | 47.10 |
| 1.464 | 62.98 | 61.05 | 45.63 |
| 1.453 | 62.64 | 60.72 | 45.38 |
| 1.442 | 60.91 | 59.05 | 44.13 |
| 1.434 | 60.66 | 58.81 | 43.95 |
| 1.426 | 59.74 | 57.91 | 43.28 |
| 1.418 | 58.81 | 57.01 | 42.61 |
| 1.401 | 57.42 | 55.66 | 41.60 |
| 1.392 | 56.40 | 54.67 | 40.86 |
| 1.384 | 55.37 | 53.68 | 40.12 |
| 1.376 | 54.74 | 53.06 | 39.66 |
| 1.369 | 54.12 | 52.45 | 39.21 |
| 1.356 | 52.45 | 50.84 | 38.00 |

| Sp. gr. | $H_3PO_4$. | $PO_4$. | $P_2O_5$. |
|---|---|---|---|
| 1.347 | 51.55 | 50.00 | 37.37 |
| 1.339 | 50.71 | 49.16 | 36.74 |
| 1.328 | 49.89 | 48.37 | 36.15 |
| 1.315 | 48.06 | 46.59 | 34.82 |
| 1.302 | 46.23 | 44.81 | 33.49 |
| 1.293 | 45.15 | 43.77 | 32.71 |
| 1.285 | 44.08 | 42.73 | 31.94 |
| 1.276 | 42.83 | 41.52 | 31.03 |
| 1.268 | 41.59 | 40.31 | 30 13 |
| 1.257 | 40.25 | 39.01 | 29.16 |
| 1.247 | 38.98 | 37.78 | 28.24 |
| 1.236 | 37.68 | 36.53 | 27.30 |
| 1.226 | 36.38 | 35.27 | 26.36 |
| 1.211 | 34.22 | 33.17 | 24.79 |
| 1.197 | 32.06 | 31.08 | 23.23 |
| 1.185 | 30.46 | 29.53 | 22.07 |
| 1.173 | 28.86 | 27.98 | 20.91 |
| 1.162 | 27.23 | 26.40 | 19.73 |
| 1.153 | 25.96 | 25.17 | 18.81 |
| 1.144 | 24.69 | 23.94 | 17.89 |
| 1.136 | 23.39 | 22.68 | 16.95 |
| 1.124 | 21.59 | 20.92 | 15.64 |
| 1.113 | 19.78 | 19.17 | 14.33 |
| 1.109 | 18.29 | 17.73 | 13.25 |
| 1.095 | 16.81 | 16.30 | 12.18 |
| 1.087 | 14.41 | 13.97 | 10.44 |
| 1.080 | 13.80 | 13.40 | 10.0 |
| 1.073 | 13 15 | 12.75 | 9.53 |
| 1.066 | 11.90 | 11.53 | 8.62 |
| 1.056 | 10.20 | 9.89 | 7.39 |
| 1.047 | 8.52 | 8.25 | 6.17 |
| 1.031 | 5.73 | 5.55 | 4.15 |
| 1.022 | 4.18 | 4.05 | 3.03 |
| 1.014 | 2.64 | 2.55 | 1.91 |
| 1.006 | 1.09 | 1.06 | 0.79 |

$H_3PO_4 = 98$; $PO_4 = 95$; $P_2O_5 \div 2 = 71$.

TABLE OF THE PERCENTAGE OF AMMONIA ($NH_4HO$), AMMONIUM ($NH_4$), AND AMMONIACAL GAS ($NH_3$) IN AQUEOUS SOLUTIONS OF AMMONIA OF DIFFERENT SPECIFIC GRAVITIES.

| Sp. gr. at 14° C. | $NH_4HO$. | $NH_4$. | $NH_3$. |
|---|---|---|---|
| 0.8844 | 74.12 | 38.118 | 36 |
| 0.8864 | 72.06 | 37.059 | 35 |
| 0.8885 | 70.0 | 36.000 | 34 |
| 0.8907 | 67.95 | 34.941 | 33 |
| 0.8910 | 66.91 | 34.410 | 32.5 |
| 0.8929 | 65.88 | 33.882 | 32 |
| 0.8953 | 63.82 | 32.824 | 31 |
| 0.8976 | 61.76 | 31.765 | 30 |
| 0.9001 | 59.71 | 30.706 | 29 |
| 0.9026 | 57.65 | 29.647 | 28 |
| 0.9052 | 55.59 | 28.588 | 27 |
| 0.9078 | 53.53 | 27.530 | 26 |
| 0.9106 | 51.47 | 26.470 | 25 |
| 0.9133 | 49.41 | 25.412 | 24 |
| 0.9162 | 47.35 | 24.353 | 23 |
| 0.9191 | 45.29 | 23.294 | 22 |
| 0.9221 | 43.23 | 22.235 | 21 |
| 0.9251 | 41.17 | 21.176 | 20 |
| 0.9283 | 39.11 | 20.118 | 19 |
| 0.9314 | 37.06 | 19.059 | 18 |
| 0.9347 | 35.00 | 18.000 | 17 |
| 0.9380 | 32.94 | 16.941 | 16 |
| 0.9414 | 30.88 | 15.882 | 15 |
| 0.9449 | 28.82 | 14.823 | 14 |
| 0.9484 | 26.76 | 13.765 | 13 |
| 0.9520 | 24.70 | 12.706 | 12 |
| 0.9556 | 22.64 | 11.647 | 11 |
| 0.9593 | 20.59 | 10.588 | 10 |
| 0.9631 | 18.53 | 9.53 | 9 |
| 0.9670 | 16.47 | 8.47 | 8 |
| 0.9709 | 14.41 | 7.41 | 7 |
| 0.9749 | 12.35 | 6.35 | 6 |
| 0.9790 | 10.29 | 5.29 | 5 |
| 0.9831 | 8.23 | 4.23 | 4 |
| 0.9873 | 6.17 | 3.18 | 3 |
| 0.9915 | 4.11 | 2.12 | 2 |
| 0.9959 | 2.06 | 1.06 | 1 |

$NH_4HO = 35$; $NH_4 = 18$; $NH_3 = 17$.

TABLE OF THE PERCENTAGE OF POTASH (KHO), OXIDE OF POTASSIUM ($K_2O$), AND POTASSIUM (K) IN AQUEOUS SOLUTIONS OF POTASH OF DIFFERENT SPECIFIC GRAVITIES.

| Specific gravity. | KHO. | $K_2O$. | K. | Boiling point. |
|---|---|---|---|---|
| 1.63 | 61.004 | 51.2 | 42.485 | 329° |
| 1.60 | 55.642 | 46.7 | 38.751 | 290 |
| 1.52 | 51.115 | 42.9 | 35.595 | 276 |
| 1.47 | 47.183 | 39.6 | 32.850 | 265 |
| 1.44 | 43.831 | 36.8 | 30.536 | 255 |
| 1.42 | 40.990 | 34.4 | 28.532 | 246 |
| 1.39 | 38.604 | 32.4 | 26.885 | 240 |
| 1.36 | 35.030 | 29.4 | 24.396 | 234 |
| 1.33 | 31.338 | 26.3 | 21 823 | 229 |
| 1.28 | 27.880 | 23.4 | 19.417 | 224 |
| 1.23 | 23.230 | 19.5 | 16.180 | 220 |
| 1.19 | 19.300 | 16.2 | 13.442 | 218 |
| 1.15 | 15.490 | 13.0 | 10.787 | 215 |
| 1.11 | 11.3 | 9.5 | 7.88 | 214 |
| ?1.06 | 5.6 | 4.7 | 3.90 | 213 |
| 1.058 | 5.84 | 4.9 | 4.07 | |

$KHO = 56$; $K_2O \div 2 = 47$; $K = 39$.

---

TABLE OF THE PERCENTAGE OF SODA (NaHO), OXIDE OF SODIUM ($Na_2O$), AND SODIUM (Na) IN AQUEOUS SOLUTIONS OF SODA OF DIFFERENT SPECIFIC GRAVITIES.

| Specific gravity. | NaHO. | $Na_2O$. | Na. | Boiling point. |
|---|---|---|---|---|
| 1.85 | 82.06 | 63.6 | 47.19 | 600° |
| 1.72 | 69.42 | 53.8 | 39.91 | 400 |
| 1.63 | 60.13 | 46.6 | 34.57 | 300 |
| 1.56 | 53.16 | 41.2 | 30.57 | 280 |
| 1.50 | 47.484 | 36.8 | 27.30 | 265 |
| 1.47 | 43.871 | 34.0 | 25.22 | 255 |
| 1.44 | 40.00 | 31.0 | 23.00 | 248 |
| 1.40 | 37.42 | 29.0 | 21.51 | 242 |
| 1.36 | 33.55 | 26.0 | 19.29 | 235 |
| 1.32 | 28.39 | 22.0 | 16.32 | 228 |
| 1.29 | 24.52 | 19.0 | 14.10 | 224 |
| 1.23 | 20.645 | 16.0 | 11.87 | 220 |
| 1.18 | 16.774 | 13.0 | 9.645 | 217 |
| 1 12 | 12.71 | 9.8 | 7.271 | 214 |
| 1.06 | 6.06 | 4.7 | 3.50 | 213 |
| 1.047 | 4.1 | 3.2 | 2.36 | |

$NaHO = 40$; $Na_2O \div 2 = 31$; $Na = 23$.

THE PROPORTION BY WEIGHT OF ABSOLUTE OR REAL ALCOHOL ($C_2H_5HO$) IN 100 PARTS OF SPIRITS OF DIFFERENT SPECIFIC GRAVITIES (FOWNES).

| Sp. gr. at 60° (15°.5 C.). | Per-centage of real alcohol. | Sp. gr. at 60° (15°.5 C.). | Per-centage of real alcohol. | Sp. gr. at 60° (15°.5 C.). | Per-centage of real alcohol. |
|---|---|---|---|---|---|
| 0.9991 | 0.5 | 0.9511 | 34 | 0.8769 | 68 |
| 0.9981 | 1 | 0.9490 | 35 | 0.8745 | 69 |
| 0.9965 | 2 | 0.9470 | 36 | 0.8721 | 70 |
| 0.9947 | 3 | 0.9453 | 37 | 0.8696 | 71 |
| 0.9930 | 4 | 0.9434 | 38 | 0.8672 | 72 |
| 0.9914 | 5 | 0.9416 | 39 | 0.8649 | 73 |
| 0.9898 | 6 | 0.9396 | 40 | 0.8625 | 74 |
| 0.9884 | 7 | 0.9376 | 41 | 0.8603 | 75 |
| 0.9869 | 8 | 0.9356 | 42 | 0.8581 | 76 |
| 0.9855 | 9 | 0.9335 | 43 | 0.8557 | 77 |
| 0.9841 | 10 | 0.9314 | 44 | 0.8533 | 78 |
| 0.9828 | 11 | 0.9293 | 45 | 0.8508 | 79 |
| 0.9815 | 12 | 0.9270 | 46 | 0.8483 | 80 |
| 0.9802 | 13 | 0.9249 | 47 | 0.8459 | 81 |
| 0.9789 | 14 | 0.9228 | 48 | 0.8434 | 82 |
| 0.9778 | 15 | 0.9206 | 49 | 0.8408 | 83 |
| 0.9766 | 16 | 0.9184 | 50 | 0.8382 | 84 |
| 0.9753 | 17 | 0.9160 | 51 | 0.8357 | 85 |
| 0.9741 | 18 | 0.9135 | 52 | 0.8331 | 86 |
| 0.9728 | 19 | 0.9113 | 53 | 0.8305 | 87 |
| 0.9716 | 20 | 0.9090 | 54 | 0.8279 | 88 |
| 0.9704 | 21 | 0.9069 | 55 | 0.8254 | 89 |
| 0.9691 | 22 | 0.9047 | 56 | 0.8228 | 90 |
| 0.9678 | 23 | 0.9025 | 57 | 0.8199 | 91 |
| 0.9665 | 24 | 0.9001 | 58 | 0.8172 | 92 |
| 0.9652 | 25 | 0.8979 | 59 | 0.8145 | 93 |
| 0.9638 | 26 | 0.8956 | 60 | 0.8118 | 94 |
| 0.9623 | 27 | 0.8932 | 61 | 0.8089 | 95 |
| 0.9609 | 28 | 0.8908 | 62 | 0.8061 | 96 |
| 0.9593 | 29 | 0.8886 | 63 | 0.8031 | 97 |
| 0.9578 | 30 | 0.8863 | 64 | 0.8001 | 98 |
| 0.9560 | 31 | 0.8840 | 65 | 0.7969 | 99 |
| 0.9544 | 32 | 0.8816 | 66 | 0.7938 | 100 |
| 0.9528 | 33 | 0.8793 | 67 | | |

## THE ELEMENTS.

| | Symbols and atomic value. | Atomic weight. |
|---|---|---|
| Aluminium ($Al_2^{VI}$) | $Al^{I}$ | 27.5 |
| Antimony ($Sb^{III}$) | $Sb^{V}$ | 122 |
| Arsenicum ($As^{III}$) | $As^{V}$ | 75 |
| Barium | $Ba^{II}$ | 137 |
| Beryllium (Glucinum) | $Be^{II}$ | 9.5 |
| Bismuth ($Bi^{III}$) | $Bi^{V}$ | 210 |
| Boron | $B^{III}$ | 11 |
| Bromine ([79.76, Stas]) | $Br^{I}$ | 80 |
| Cadmium | $Cd^{II}$ | 112 |
| Cæsium | $Cs^{I}$ | 133 |
| Calcium | $Ca^{II}$ | 40 |
| Carbon ($C^{II}$) | $C^{IV}$ | 12 |
| Cerium ($Ce^{II}$) ([136, Mendelejeff]) | $Ce^{VI}$ | 92 |
| Chlorine ([35.368, Stas]) | $Cl^{I}$ | 35.5 |
| Chromium ($Cr_2^{VI}$) | $Cr^{VI}$ | 52.5 |
| Cobalt ($Co^{II}$) | $Co^{VI}$ | 58.8 |
| Copper | $Cu^{II}$ | 63.5 |
| Davyum? | | |
| Decipium? | $Dp^{II}$ | |
| Didymium ([138, Mendelejeff]) | $D^{II}$ | 96 |
| Erbium? ([171, Mendelejeff]) | $Eb^{II}$ | 112.6 |
| Fluorine | $F^{I}$ | 19 |
| Gallium | $Ga^{IV}$ | 69.8 |
| Glucinum. *See* Beryllium. | | |
| Gold ($Au^{I}$) | $Au^{III}$ | 196.7 |
| Hydrogen | $H^{I}$ | 1 |
| Indium ([113, Mendelejeff]) | $In^{VI}$ | 75.6 |
| Iodine ([126.538, Stas]) | $I^{I}$ | 127 |
| Ilmenium? | $Il^{V}$ | |
| Iridium | $Ir^{IV}$ | 197 |
| Iron ($Fe^{II}$ & $Fe_2^{VI}$) | $Fe^{VI}$ | 56 |
| Lanthanum ([139, Mendelejeff]) | $L^{II}$ | 92 |
| Lavoisium? | | |
| Lead ($^{II}Pb^{II}$) | $Pb^{IV}$ | 207 |
| Lithium ([7.004, Stas]) | $Li^{I}$ | 7 |
| Magnesium | $Mg^{II}$ | 24 |
| Manganese ($Mn^{II}$ & $Mn^{IV}$) | $Mn^{VI}$ | 55 |
| Mercury | $Hg^{II}$ | 200 |
| Molybdenum | $Mo^{VI}$ | 96 |
| Neptunium | $Np^{V}$ | |
| Nickel ($Ni^{II}$) | $Ni^{VI}$ | 58.8 |

| | Symbols and atomic value. | Atomic weight. |
|---|---|---|
| Niobium | $Nb^{V}$ | 97.6 |
| Nitrogen ($N^{I}$ & $N^{III}$) (14.009, Stas) | $N^{V}$ | 14 |
| Osmium | $Os^{IV}$ | 199 |
| Oxygen (15.96, Stas) | $O^{II}$ | 16 |
| Pallabium | $Pd^{IV}$ | 106.5 |
| Philippium? | $Pp^{II}$ | |
| Phosphorus ($P^{III}$) | $P^{V}$ | 31 |
| Platinum (197.86, Andrews) | $Pt^{IV}$ | 198 |
| Potassium (39.04, Stas) | $K^{I}$ | 39 |
| Rhodium | $Rh^{IV}$ | 104.3 |
| Rubidium | $Rb^{I}$ | 85.3 |
| Ruthenium | $Ru^{IV}$ | 104.2 |
| Selenium or Selenion | $Se^{VI}$ | 79.5 |
| Silicon | $Si^{IV}$ | 28 |
| Silver (107.66, Stas) | $Ag^{I}$ | 108 |
| Sodium (22.96, Stas) | $Na^{I}$ | 23 |
| Strontium | $Sr^{II}$ | 87.5 |
| Sulphur ($S^{II}$ & $S^{IV}$) | $S^{VI}$ | 32 |
| Tantalum | $Ta^{V}$ | 182 |
| Tellurium | $Te^{VI}$ | 129 |
| Terbium? | $Tb^{II}$ | |
| Thallium (203, Crookes) | $Tl^{III}$ | 204 |
| Thorinum or Thorium | $Th^{II}$ | 232 |
| Tin ($Sn^{II}$) | $Sn^{IV}$ | 118 |
| Titanium | $Ti^{IV}$ | 50 |
| Tungsten | $W^{VI}$ | 184 |
| Uranium (240, Mendelejeff) | $U^{VI}$ | 120 |
| Vanadium | $V^{V}$ | 51.3 |
| Ytterbium? | Yb | |
| Yttrium (90, Mendelejeff) | $Y^{II}$ | 61.7 |
| Zinc | $Zn^{II}$ | 65 |
| Zirconium | $Zi^{IV}$ | 89.5 |
| Total . . 72 | | |

*The quantivalence* or atomic value of some elements is, apparently, variable; in the above Table the full coefficients are given in the column of symbols, other common values in brackets.

*Atomic weights* are sometimes obscurely termed *equivalents*.

# INDEX.


*Abies balsamea*, 472
*canadensis*, 472
*excelsa*, 470
*Abrus precatorius*, 418
Absolute alcohol, 425, 515
*Acacia catechu*, 357
*Acacia suma*, 357
*Acaciæ gummi*, 114
*gummi*, impurities in, 615
Acetate of ammonium, solution of, 92
amyl, 441
copper, 192
ethyl, 300
iron, 149
lead, 210
morphia, 382
potassium, 69
sodium, 82
zinc, 133
**Acetates**, 297
analytical reactions of, 300
decomposition of aqueous solution of, 298
volumetric estimation of, 554
Acetic acid, 299, 460
ether, 300
glacial, 299, 517
volumetric estimation of free, 554
Acetone, 300
Acetonitrate of barium, 126
of iron, 157
*Acetum*, 298
*cantharidis*, 298
*colchici*, 298
*destillatum*, 298
impurities in, 300, 615
*lobeliæ*, 298
*opii*, 298
*sanguinariæ*, 298
*scillæ*, 298
Acetyl, 298
Acid, acetic, 299, 460
acetic, glacial, 299, 517
aconitic, 325
amyric, 471
angelic, 463
arabic, 405
arachidic, 460
arsenic, 169
arsenious, 168
benzoic, 335, 473, 616
boracic, 333
bromic, 296
butyric, 361, 460
camphoretic, 467
camphoric, 467
cantharidic, 468
caproic, 460
caprylic, 460
carbazotic, 445
carbolic, 444, 515
carbolic, impure, 444
carbonic, 30, 312
carminic, 337
catechuic, 357
cathartic, 415
cathartogenic, 415
cerotic, 460
cetraric, 337
chavicic, 398
chloric, 292, 294
cholalic, 457
chromic, 238
chrysammic, 446
chrysophanic, 337
cinnamic, 336, 473
citric, 324
colopholic, 469
colophonic, 469
copaivic, 471
cresotic, 445
cresylic, 444, 445

Acid—
cryptophanic, 494
cuminic, 465
cyanic, 337
dextroracemic, 320
dextrotartaric, 320
dithionic, 345
equisitic, 325
erucic, 459
ethyl sulphuric, 429
eugenic, 464
ferulaic, 472
filicic, 459
fluoric, 341
formic, 338, 460
gallic, 338, 357
gambogic, 472
gaultheric, 442
gelseminic, 396
gentianic, 418
gentisic, 418
glacial acetic, 299, 515
guaiaconic, 418
guaiaretic, 418
guaiaretinic, 418
gummic, 405
hemidesmic, 338
hippuric, 338, 497, 500
hydriodic, 273
hydrobromic, 270
hydrochloric, 29, 266
common, 266
dilute, 266
hydrocyanic, 279, 280
dilute, 280
hydroferridcyanic, 340
hydroferrocyanic, 339
hydrofluoric, 341
hydrosulphuric, 301
hypochlorous, 292
hypophosphoric, 351
hypophosphorous, 342
hyposulphurous, 344
iodic, 296
jalapic, 419
lactic, 346
lævoracemic, 320
lævotartaric, 320
larixinic, 357
lauric, 460
lithic, 359
lupulinic, 472
malic, 347
mastichic, 470
Acid—
meconic, 348, 489
melissic, 460
metaboracic, 334
metagummic, 405
metantimonic, 181
metaphosphoric, 348
metastannic, 243
mimotannic, 357
mucic, 413
muriatic, 266
myristic, 460
myrrhic, 473
naphthalic, 336
nitric, 285, 287
dilute, 288
nitrohydrochloric, 186, 288
dilute, 288
nitromuriatic, 288
nitrous, 349
œnanthylic, 460
oleic, 459
ophelic, 350
opianic, 382
orthophosphoric, 350
oxalic, 316
chemically pure, 316
palmitic, 458
paratartaric, 320
parietinic, 337
pelargonic, 460
pentathionic, 345
perchloric, 294
phenic, 444
phosphomolybdic, 492
phosphoric, 25, 329, 350
dilute, 329
glacial, 330
phosphorous, 350
phthalic, 336
picric, 445
pimaric, 469
pinic, 469
propionic, 460
prussic, 279
pyrogallic, 358
pyroligneous, 297
pyrophosphoric, 351
racemic, 320
rheic, 337
rhubarbaric, 337
rutic, 459
saccharic, 413
salicylic, 442, 445

Acid—
salicylous, 442
santonic, 420
sarcolactic, 347
sclerotic, 470
sclerotinic, 470
silicic, 352
stannic, 244
stearic, 456, 460
succinic, 354
sulphethylic, 424
sulphindigotic, 290
sulphindylic, 290
sulphocarbolic, 445
sulphocyanic, 354
sulphophenic, 445
sulphosalicylic, 446
sulphovinic, 424
sulphuric, 308
sulphuric, aromatic, 310
dilute, 310
sulphurous, 305
sulphydric, 301
sylvic, 469
tannic, 355
tartaric, 318, 319
tetrathionic, 345
tiglic, 459
trinitrocarbolic, 445
trithionic, 345
tropic, 393
uric, 358
valerianic, 359, 460
Acid carbonate of potassium, 71, 314
carbonate of sodium, 83
salts, 72, 302
solution of arsenic, 172
tartrate of potassium, 61, 78
tartrate of sodium, 79
Acidimetry, 552
Acids, analytical detection of, 366
antidotes to, 265
definition of, 263
free, estimated, 585
of chlorine, 292
quantitative estimation of, 554
volumetric estimation of official, 554
Acidulous radicals, formulæ and quantivalence of, 66, 122, 264
radicals, qualitative detection of, 362
Acidulous—
radicals, quantitative estimation of salts of, 585
radicals, tables to aid in the detection of, 365, 366
*Acidum aceticum*, 299
*aceticum*, impurities in, 615
*aceticum dilutum*, 299
*aceticum glaciale*, 299
*aceticum glaciale*, impurities in, 615
*arseniosum*, 168
*benzoicum*, 335
*boracicum*, impurities in, 615
*carbolicum*, 444
*carbolicum impurum*, 444
*chromicum*, 238
*citricum*, 324
*citricum*, impurities in, 615
*gallicum*, 357
*gallicum*, impurities in, 615
*hydrochloricum*, 29, 266
*hydrochloricum*, impurities in, 615
*hydrochloricum dilutum*, 266
*hydrocyanicum dilutum*, 280
*hydrocyanicum dilutum*, impurities in, 615
*lacticum*, 346
*nitricum*, 287
*nitricum*, impurities in, 615
*nitricum dilutum*, 288
*nitro-hydrochloricum dilutum*, 288
*nitro-muriaticum*, 288
*oxalicum*, 316
*oxalicum*, impurities in, 615
*phosphoricum dilutum*, 329
*phosphoricum dilutum*, impurities in, 615
*sulphuricum*, 310
*sulphuricum*, impurities in, 616
*sulphuricum aromaticum*, 310
*sulphuricum dilutum*, 310
*sulphurosum*, 306
*sulphurosum*, impurities in, 616
*tannicum*, 355
*tannicum*, impurities in, 616
*tartaricum*, 319
*tartaricum*, impurities in, 616
*valerianicum*, 361
*Acipenser*, 453
*Aconiti ferocis radix*, 392
*Aconiti folia*, 392

*Aconiti heterophylli radix*, 393
*radix*, 392
*Aconitia*, 392
impurities in, 616
Aconitic acid, 325
Aconitina, 392
Aconitine, 392
*Aconitum ferox*, 392
*heterophyllum*, 393
*napellus*, 392
Acorin, 406
*Acorus calamus*, 466
Acrinyl sulphocyanate, 447
Acrolein, 448
*Actea racemosa*, 400
*Adeps benzoatus*, 458
*præparatus*, 458
*præparatus*, impurities in, 616
Adhesion, 56
Adipocire, 453
*Ægle marmelos*, 357
Aerated bread, 411
water, 85
Æsculin, 396
*Æther*, 427
*aceticus*, 431
*aceticus*, impurities in, 616
*fortior*, 430
*fortior*, impurities in, 616
*purus*, 430
*purus*, impurities in, 616
Affinity, chemical, 38
Agate, 352
Air, composition, 26
influence of animals and plants on, 19
nitrogen in the, 26
oxygen in the, 15
relative weight of the, 26
weight of 1 cubic cent., 537
weight of 100 cub. inches, 537
Ajwain oil, 463
Alabaster, 105
Albumen, 449
detection of, in urine, 494
vegetable, 453
*Albumen ovi*, 449
Albumenoid substances, 450
Alchemy, 13
*Alcohol*, 422
absolute, 425, 515
amylic, 441, 515
and allied bodies, 422 *et seq.*
benzylic, 473
*Alcohol*—
butylic, 361
cinnamic, 473
*dilutum*, 424
ethylic, 424
*fortius*, 423
from sugar, 410
glyceric, 448
in 100 parts of spirits of different densities, Table showing the proportion of, 634
methylic, 434
phenic, 444
quantitative estimation of, 610
radicals, 433
real, 425
test for impurities in, 616
test for purity of, 425
*Alcyhol amylicum*, 441
*amylicum*, impurities in, 616
Alcoholates of chloral, 440
Alcoholometer, 533
Alcohols, 433
Aldehyd, 426
benzoic, 336
euodic, 466
lauric, 466
rutic, 466
Aldehyds, 434
aromatic, 466
Ale, 405, 423
Alexandrian senna, 415
Algaroth's powder, 181
Alizarin, 476
Alkalies, analytical separation of the, 100
antidotes to, 265
quantitative estimation of the, 544
Alkalimetry, 551
Alkaline carbonates, volumetric estimation of the, 544
earths, 126
solution of arsenic, 168
**Alkaloids**, 379
antidotes to the, 380
distinguished, 380, 392
nomenclature of, 380
poisonous, examination for, 490 *et seq.*
Alkanet, 475
*Alkanna tinctoria*, 475
Allotropes, 406

Allotropic bodies, 406
Allotropy, 406
Alloxan, 359
Alloy, 196
Alloys, analysis of, 374
Allyl, 448
cyanide, 448
sulphide of, 448
sulphocyanate of, 448
Almond-oil, 459
Almonds, oil of bitter, 414, 443
oil of bitter, test for nitrobenzol in, 415
*Aloe barbadensis*, 446
*socotrina*, 446
Aloes, 446
Aloins, 446
*Alstonia scholaris*, 396
Alum cake, 138
Alum, 137
chrome-, 137
dried, 138
-flour, 137
iron-, 138
potash-, 138
roche or rock, 138
root, 357
soda-, 137
*Alumen*, 137
impurities in, 616
*exsiccatum*, 138
Alumina, 139
**Aluminium**, 136
analytical reactions of, 139
and ammonium, sulphate of, 137
and sodium, double chloride of, 137
bronze, 137
derivation of word, 32
detection of, in presence of iron and zinc, 162
hydrate of, 139
oxide of, 139
quantitative estimation of, 577
separation of, from chromium and iron, 240
silicate of, 137
sulphate of, 138
Amalgam, 196
ammonium, 90
Amber, 354
oil of, 354
American senna, 415
turpentine, 466
Amianth, 352
Amide, 205
Amido-chloride of mercury, 205
Amidogen, 205
Amines, 379
Ammonia, 91
detected by Nessler test, 548
fetid spirit of, 93
gas, composition of, 89
gas in solutions of ammonia of different densities, Tables showing the amount of, 632
in drinking waters, 548
preparation of, 91
solution of, 91
type, 377
volcanic, 90
volumetric estimation of solutions of, 547
Ammoniacal liquor, 90
salts, sources of, 90
Ammoniacum, 472
*Ammoniæ* (*vide Ammonii*).
Ammoniated mercury, 205
mercury, varieties of, 206
*Ammonii acetatis, liquor*, 92
*aromaticus, spiritus*, 93
*benzoas*, 94, 336
*benzoas*, impurities in, 616
*bromidum*, 94
*carbonas*, 92
*carbonas*, impurities in, 616
*chloridum*, 90
*chloridum*, impurities in, 616
*citratis, liquor*, 94
*fœtidus, spiritus*, 93
*fortior, liquor*, 91
*iodidum*, 94
*liquor*, 91
*nitras*, 93
*nitras*, impurities in, 616
*phosphas*, 94
*sulphas*, 90
Ammonio-chloride of mercury, 196
-citrate of iron, 153
-ferric sulphate, 138
-magnesian phosphate, 120, 330
-nitrate of silver, 177, 206
-sulphate of copper, 177, 206
-tartrate of iron, 155

**Ammonium**, 89
acetate, 92
-amalgam, 90
analytical reaction, 98
and magnesium, arseniate of, 120
and magnesium phosphate, 120
and platinum, double chloride, 98, 248
arseniate, 170
benzoate, 94, 336
bicarbonate, 93
bromide, 94, 271
carbamate, 93
carbonate, 92
carbonate, solution of, 93
chloride, 90
citrate, 94
cyanate, 337
derivation of word, 32
derivatives, 206
hydrate, 91
iodide, 273
molybdate, 492
nitrate, 93
oxalate, 95
periodide, 274
phosphate, 94
potassium, and sodium, separation of, 100
quantitative estimation of, 572
salts, source of, 90
salts, volatility of, 99
sulphate, 90
sulphide, 96
sulphydrate, 96
tartrate of, 98
volumetric estimation of carbonate of, 544
Amorphous, meaning of, 184
phosphorus, 328
*Amygdala amara*, 414
*dulcis*, 414
Amygdalin, 414
Amyl, 441
acetate, 441
nitrite, 442
valerianate, 441
Amylaceous substances, 401
Amylamine, 380
Amylic alcohol, 441
*Amylum*, 401
impurities in, 616
Amyric acid, 471
Amyrin, 471
*Anacyclus pyrethrum*, 470
Analogies between chlorine, bromine, and iodine, 278
of sodium and potassium salts, 87
Analogy of carbon, boron, and silicon, 333, 354
of oxygen, sulphur, selenium, and tellurium, 303
of salts, 87
Analysis, 100
aided by sifting, 370
blowpipe, 371
gas-, 358, 481
gravimetric, 511
meaning of word, 60
organic, 597
practical, 99
proximate, 597
qualitative, 99
quantitative, 510
spectral, 482
systematic, for the detection and separation of the metals, 100, 125, 164, 186, 189, 223, 373
ultimate, 597
volumetric, 511, 543
Analysis and synthesis, 60
of gases and vapors, 481
of insoluble salts, 372 *et seq.*
of medicines, 378
of salts, 369
of substances having unknown properties, 372
Analytical chemists, 14, note.
detection of the acidulous radicals of salts soluble in water, 362
memoranda, 260, 265
**Analytical reactions of—**
acetates, 300
albumen, 449
alcohol, 425
aldehyd, 426
aluminium, 139
ammonium, 98
amygdalin, 414
antimony, 184, 484
arsenicum, 171, 484
atropia, 393
barium, 103

**Analytical reactions of—**
beberia, 394
benzoates, 336
berberia, 394
bile, 457
bismuth, 254
borates, 334
bromides, 272
brucine, 391
cadmium, 249
caffeia, 398
calcium, 114
carbonates, 314
chloral hydrate, 439
chlorates, 295
chlorides, 269, 486
chromates, 238
chromium, 240
citrates, 326
cobalt, 234
conia, 395
copper, 192
cyanides, 282
ferric salts, 160
ferridcyanides, 341
ferrocyanides, 339
ferrous salts, 159
fluorides, 341
formates, 338
gallic acid, 357
glucosides, 413
glycerine, 448
gold, 246
guaiacin, 418
hippurates, 339
hydrocyanic acid, 282, 487
hypochlorites, 293
hypophosphites, 344
hyposulphites, 345
iodides, 274
iron, 159
lactates, 347
lead, 213, 485
lithates, 359
lithium, 227
magnesium, 120
malates, 347
manganese, 232
meconates, 348, 489
mercuric salts, 205, 207
mercurous salts, 204, 207
metaphosphates, 348
morphia, 381, 489
nickel, 235

**Analytical reactions of—**
nicotia, 397
nitrates, 288
nitrites, 349
nitrous ether, 430
oxalates, 317, 486
phosphates, 330
phosphites, 350
platinum, 247
potassium, 77
pyrogallic acid, 358
pyrophosphates, 348
quinia, 386
salicin, 420
silicates, 353
silver, 218
sodium, 88
starch, 401
strontium, 228
strychnia, 390, 488
succinates, 354
sugar, 409
sulphates, 311
sulphides, 304
sulphites, 307
sulphocyanates, 354
sulphuric acid, 311
tannic acid, 356
tartrates, 322
theia, 398
tin, 243
urates, 359
veratia, 399
zinc, 135
*Anamirta cocculus*, 420
*Anchusa tinctoria*, 475
Anchusin, 475
*Andrographis paniculata*, 400
*Andropogon citratus*, 467
*nardus*, 464
*schœnanthus*, 465
Aneroid barometer, 512
Anethene, 463
Anethol, 463
Angelic acid, 463
powder, 181
Angelica, 463
Angustura bark, 400
Anhydride, acetic, 299
antimonic, 181
antimonius, 181
boracic, 333
carbonic, 313
chlorochromic, 239

Anhydride—
chromic, 238
nitric, 285, 288
nitrous, 289
phosphoric, 25, 350
silicic, 353
sulphuric, 310
sulphurous, 305
Anhydrides, 84
Anhydrous bodies, 84
nitric acid, 287
perchloride of iron, 146
sulphate of copper, 192
Aniline, 445, 478
colors, 478
Animal charcoal, 112
decolorizing power of, 112
rouge, 476
Animals and plants, complementary, action of air, 19
Anise, 463
Aniseed-oil, 463
Anise-fruit, 463
Annatto, 476
*Anthemidis flores*, 400, 463
*Anthemis nobilis*, 463
Anthracene, 446
Anthracite, 241
Antichlor, 306
**Antidotes** to acids, 265
alkalies, 265
alkaloids, 380
antimony, 185
arsenic, 177
barium, 103
carbolic acid, 444
copper, 193
cyanides, 283
hydrochloric acid, 269
hydrocyanic acid, 283
lead, 214
mercury, 208
nitric acid, 291
oxalic acid, 317
silver, 219
sulphuric acid, 312
tin, 244
zinc, 136
Antimonial wine, 182
Antimoniate of potassium, 88
sodium, 88
Antimonic anhydride, 181
chloride, 180
oxide, 181
*Antimonii chloridi, liquor*, 180, 580
*et potassæ tartras*, 182
*oxidum*, 181
*oxidum*, impurities in, 616
*oxysulphuretum*, 182
*sulphuretum*, 182
Antimonious anhydride, 181
chloride, 180
oxide, 181
oxychloride, 180
*Antimonium nigrum*, 179
*nigrum*, impurities in, 616
*sulphuratum*, 182
*tartaratum*, 182, 319, 373
*tartaratum*, quantitative estimation of antimony in, 373
Antimoniuretted hydrogen, 185
**Antimony**, 166, 179, 319, 517
analytical reactions of, 184
and arsenic, analytical separation of, 186
and potassium, tartrate of, 181, 319
antidote to, 185
black, 179
butter of, 180
chloride of, 180
crude, 179
derivation of word, 32
from arsenic, to distinguish, 175
hydride of, 185
in organic mixtures, detection of, 485
oxide of, 181
oxychloride of, 180
oxysulphide of, 182
pentachloride of, 180
potassio-tartrate of, 181
quantitative estimation of, 579
solution of chloride of, 180
sulphide of, 179, 184
sulphur salts of, 182
sulphurated, 182
tartarated, 181
Antiseptic, 444
Antozone, 238, 275, 502
Apomorphia, 383
Aporetine, 337
Apothecaries, 14
Apparatus, xi.
for experiments, xi.

Apparatus—
for gravimetric analysis, 568, 576, 581, 582, 587, 591, 597, 598, 599, 600
for volumetric analysis, 543
list of, xi.
Apple, essence of, 441
oil, 441
wine, 423
*Aqua*, 126
*Aqua acidi carbolici*, 444
*acidi carbonici*, 312
*ammoniæ*, 91
*ammoniæ fortior*, 91
*amygdalæ amaræ*, 414
*anethi*, 462
*aurantii floris*, 462
*aurantii floris*, impurities in, 616
*camphoræ*, 467
*chlorinii*, 28, 268
*chloroformi*, 438
*creasoti*, 444
*destillata*, 127
*destillata*, impurities in, 616
*fortis*, 288
*laurocerasi*, 414
*menthæ piperitæ*, 462
*viridis*, 462
*regia*, 186, 288
*rosæ*, 462, 465
Arabin, 114
Arachidic acid, 460
Arachis hypogeia, 459
Arachis oil, 459
Araroba powder, 337
*Arbor Dianæ*, 218
Arbutin, 357, 415
Archil, 477
*Arctostaphylos uva ursi*, 415
Are, 522
*Areca catechu*, 357
Areca nuts, 357
Argal, 318
Argent-ammon-ammonium, nitrate of, 206
*Argenti cyanidum*, 219
*nitras*, 218
*nitras*, impurities in, 616
*oxidum*, 218
*oxidum*, impurities in, 616
Argentic chloride, sulphide, etc., *vide* salts of silver.
Argentiferous galena, 216
*Argentum*, 33, 217
*Argentum purificatum*, 217
*purificatum*, impurities in, 616
Argol, 318
Armenian bole, 476
*Armoraciæ radix*, 463
Arnatto, 476
Arnica, 469
*Arnicæ radix*, 469
Arnicin, 469
Arnotto, 476
Aromatic aldehydes, 464
Arrowroot-starch (fig.), 403
Arseniate of ammonium, 169
barium, 104, 177
calcium, 177
copper, 177
iron, 170, 562
magnesium and ammonium, 120
silver, 177
sodium, 169
volumetric estimation of, 556
zinc, 177
**Arseniates**, 169
Arsenic acid, 169
and arsenical solutions, volumetric estimation of official, 559
anhydride, 169
in carbonate of potassium, solution of, 167
in hydrochloric acid, solution of, 168
odor of, 168
white, 167
**Arsenical ores**, 167
sulphur, 176
*Arsenici iodidum*, 166
**Arsenicum**, 167, 168
analytical reactions of, 171
antidotes to, 177
and antimony, analytical separation of, 186
bromide, 166
chloride, 166
derivation of word, 32
detection of, in metallic copper, 173
detection of, in organic mixtures, 484
Fleitmann's test for, 174

**Arsenicum**—
from antimony, to distinguish, 175
hydride of, 174
iodide of, 166
Marsh's test for, 173
quantitative estimation of, 560, 579
red native sulphide of, 167
reduction of arseniates to arsenites, 169
Reinsch's test for, 172
sources of, 167
sulphide of, 167, 176
yellow native sulphide of, 167
Arsenide of cobalt, 234
Arsenio-sulphide of iron, 167
of nickel, 235
Arsenious acid, 168
anhydride, 168
**Arsenites**, 168
Arsenite of copper, 177
potassium, 167
silver, 177
sodium, 168
Arseniuretted hydrogen, 174
Art of chemistry, 13
Artemisia, 421
Artificial alkaloids, 379
gastric juice, 454
*Asafœtida*, 472
Asbestos, 352
Asclepedin, 400
*Asclepias tuberosa*, 400
Ash, 101
black-, 87
bone-, 110
soda-, 87
Ashes, analysis of (mixed solids), 372
Asparagin, 347
Aspartate of ammonium, 347
Aspirator, water, 309
-ate, meaning of, 72
Atees, 393
Atis, 393
Atmosphere, carbonic acid in, 313
composition of, 26
nitrogen in, 26
oxygen in, 15, 26
Atmospheric pressure, measurement of, 511
Atom, definition of, 57
weights, definition of, 51
Atomic proportions, 197, 200
theory, 50
weights, 51
relation of specific heat to, 539
Atomicity, 53
Atoms, 50, 51
quantivalence of, 55
definition of, 58
*Atropa belladonna*, 393
Atropia, tropate of, 393
*Atropia*, 393
impurities in, 616
*Atropiæ sulphas*, 393
*sulphas*, impurities in, 616
Atropine, 393
Attar of rose, 465
*Aurantii cortex*, 400, 463
*amari cortex*, 463
*dulcis cortex*, 463
*flores*, 463
fructus, 463
Auric chloride, 245
sulphide, 246
*Aurum*, 34
*Avenæ farina*, 401
Avignon grains, 475
Avogadro's and Ampère's law, 52, 53
*Azadiracta indica*, 400

Bael fruit, 357
Bahia powder, 337
Baking-powder, 410
Balance, 519
Balloons, coal-gas for, 21
hydrogen for, 21
Balm-of-Gilead fir, 471
Balsam, Canada, 471
copaiva, 471
fir, 466
Gurjun, 471
Peru, 473
Storax, 473
Tolu, 473
*Balsamodendron myrrha*, 473
*Balsamum Peruvianum*, 473
*Peruvianum*, impurities in, 616
*Tolutanum*, 473
Balsams, 473
*Baptisia tinctoria*, 400
Barbadoes aloes, 446
Barbaloin, 446

Barberry, 394
Barff's protected iron, 141
Baric chloride, nitrate, etc., *vide* salts of barium.
*Barii carbonas*, 102
**Barium**, 102
acetonitrate of, 126
analytical reactions of, 103
and calcium, separation of, from magnesium, 124
antidotes to, 103
arseniate of, 104
carbonate of, 104
native, 102
chloride of, 102
chromate of, 103
derivation of word, 32
detection of, in presence of calcium and magnesium, 124
hydrate of, 103
nitrate of, 102
peroxide of, 103
phosphate of, 104, 332
oxalate of, 104, 317
quantitative estimation, 573
salts, antidote to, 103
silico-fluoride of, 104
sulphate of, 103
sulphide of, 102
sulphite, 307
Barley-starch (fig.), 403
Barley-starch, 412
Barometer, 511
Baryta, 102
-water, 103
Basalt, 137
Base, meaning of, 263
organic, 379
Bassorin, 114, 405
Bastard saffron, 476
Basylous radicals, 122
Bath brick, 352
Bauxite, 137
Bay rum, 423
Bay-salt, 81
Bearberry, 357
Beaver, 469
Beberia or beberine, 393
*Beberiæ sulphas*, 394
*sulphas*, impurities in, 616
Bebirine, 393
Beer, 405
Beeswax, 458
Beetroot, 410
*Belæ fructus*, 357
*Belladonnæ folia*, 393
*radix*, 393
Bell-metal, 241
Bend glass tubes, to, 16
Benne oil, 460
Benzene, 443
Benzine-collas, 443
Benzoate of ammonium, 94, 336
Benzoated lard, 458
Benzoates, 335
Benzoic acid, 335, 515, 517
aldehyde, 336
glycocine, 338
Benzoin, 335, 473
*Benzoinum*, 335, 473
Benzol, 443
Benzyl, hydrate of, 336
hydride of, 414, 473
Benzyl, benzoate of, 473
cinnamate of, 473
Benzylic alcohol, 473
Berberia or berberine, 394
*Berberis cortex*, 394
Bergamot oil, 463
Berlin blue, 477
red, 476
Berthollet's laws, 378
Beryllium, 635
Betel nuts, 357
Bi-, the prefix, 72
Bibasic, *vide* Dibasic.
Bibirine, 393
Bibulous paper, 108
Bicarbonate of ammonium, 92
of potassium, 71, 321, 323, 547
of potassium, chemically pure, 550
of sodium, 83, 321, 323, 547
of sodium, chemically pure, 550
Bichromate of potassium, 239
Bikh, 392
Bile, 457
detection of, in urine, 495
test for presence of, 457
Biliary calculi, 506
Bish, 392
**Bismuth**, 250, 517, 534, 580
and ammonium, solution of citrate of, 253
analytical reactions of, 254

**Bismuth—**
carbonate of, 253
citrate of, 253
derivation of word, 34
hydrate, 254
-lozenge, 252
nitrate, 250
oxide, 252
oxy-salts of, 252
quantitative estimation of, 580
-salts, composition of, 253
subcarbonate or oxycarbonate, 252
subnitrate or oxynitrate, 252
sulphate, 253
sulphide, 254
*Bismuthi carbonas*, 252
*carbonas*, estimation of bismuth in, 580
*carbonas*, impurities in, 616
*Bismuthi oxidum*, 252
*oxidum*, impurities in, 617
*subcarbonas*, 252
*subnitras*, 251
*Bismuthum*, 251
*purificatum*, 251
*purificatum*, impurities in, 617
Bisulphide of carbon, 433
Bisulphite of lime, 306
Bitter almonds, oil of, 414, 459
cassava, 401
principles, 400
-sweet, 398
Bittern, 270
Bituminous coal, 241
Bivalence, 55
Bivalent radicals, 56, 122
*Bixa orellana*, 476
Bixin, 476
Black antimony, 179
ash, 87
-band, 140
bone-, 111
cherry bark, 415
coloring matters, 477
dyes, 477
flux, 169
hydrate of iron, 156
ink, 356
-lead, 30
oak, 475
oxide of copper, 193
oxide of iron, 156
Black—
oxide of manganese, 230
oxide of mercury, 204
pepper, 397
Bladder-green, 477
*Blanc de Perle*, 252
Bleaching by chlorine, 29
-liquor, 114
-powder, 113
Blende, 130
Block tin, 241
Blood, 451, 501
detection of, in organic matter, 501
hydrocyanic acid in the, 284
Blood-root, 398
Blood-root vinegar, 298
Blood stains, 501
Blowpipe-analysis, 371
Blue cohosh, 400
coloring-matters, 477
copperas, 143
flag, 400
indigo, 291
ointment, 195
pill, 195
Prussian, 160, 340
stone, 192
Turnbull's, 341
vitriol, 143, 192
Boiled oil, 459
Boiling-points of various substances, 515
-point, definition of, 515
Boldo, 464
"Bonds" (Frankland), 137
Bondue seeds, 400
Bone-ash, 110
-black, 111
-earth, 327
Bones, composition of, 327
Boracic acid, 333
anhydride, 333
as an antiseptic, 334
Borates, 333
analytical reactions of, 334
Borax, 333
volumetric estimation of, 546
Borax bead, 232
Bordeaux turpentine, 466
Boric acid (see Boracic acid), 333
Borneène, 467
Borneo camphor, 467

**Boron**, 333
chloride of, 333
derivation of word, 333
flame, 334
fluoride of, 333
Borotartrate of potassium, 334
*Bos taurus*, 457
*Boswellia*, 473
Bourdon barometer, 512
Boyle's law, 52
Brandy, 423, 426
Brass, 130
Brazil powder, 337
Brazil wood, 476
Bread, 401, 410
aerated, 411
making, 410
Breidin, 471
Brezilin, 476
Bright's disease, 494
Britannia metal, 179, 209
British gum, 404
Bromal, 440
alcoholates of, 440
hydrate of, 440
Bromate of potassium, 76
Bromates, 271, 296
Bromic acid, 76, 296
Bromide of ammonium, 271
arsenicum, 166
ethyl, 432
iron, 146
potassium, 76, 271
potassium, volumetrical estimation of, 556
of silver, 219
of sulphur, 305
Bromides, 270
analytical reactions of, 272
quantitative analysis of, 556, 586
separation of, from chlorides and iodides, 276
**Bromine**, 270
analytical separation of, 272
derivation of word, 33
its analogy to chlorine and iodine, 273
solution of, 271
volumetric estimation of free, 586
*Bromum*, 270
*Brominium*, 270
impurities in, 617
Bronze, 241
aluminium, 137
coinage, 191
Bronzing-powder, 244
Broom-tops, 398
Brown coloring matters, 477
hæmatite, 140
rosin, 469
sugar, 409
Brucia, 391
Brucine, 391
Brunswick green, 177
Bryoidin, 471
*Buchu folia*, 464
Buchu, oil of, 464
Buckthorn green, 477
-juice, 416
"Bumping," 281
Bunsen gas-burners, 23
valve, 589
Burette, Mohr's, 543
Burgundy pitch, 470
Burners, gas, 17, 23
Burnett's disinfecting fluid, 131
Burnt ochre, 476
sugar, 412
umber, 477
*Butea frondosa*, 357
Butter, 451
of antimony, 180
of cacao, 458
of cocoa, 458
of kokum, 458
of orris, 465
Butyl, 361
chloral, 440
sulphocyanate, 434
Butylic alcohol, 361
Butyrates, 361
Butyric acid, 361, 434, 460
Buxine, 394
*Buxus sempervirens*, 394
By-products, 200

Cacao butter, 458
Cacaotine, 458
*Cadmii iodidum*, 249
*iodidum*, impurities in, 617
**Cadmium**, 249
analytical reactions of, 249
carbonate, 249
derivation of word, 34
hydrate of, 250
iodide of, 249

**Cadmium—**
nitrate, 249
sulphate, 249
sulphide of, 249
*Cæsalpinia braziliensis*, 476
*bonducella*, 400
Cæsium, 635
Caffeia, 398
Cajuput oil, 464
Cajuputene, 464
Cajuputol, 464
Caking coal, 241
Calabar bean, 397
Calamine, 130
Calamus draco, 470
Calcic sulphate, phosphate, etc., *vide* salts of calcium.
*Calcii carbonas præcipitata*, 107
*carbonas præcipitata*, impurities in, 617
*chloratæ, liquor*, 114
*chloridum*, 105
*chloridum*, impurities in, 617
*hydras*, 107
*hypophosphis*, 343
*phosphas*, 110
*phosphas*, impurities in, 617
*saccharatus, liquor*, 107
Calcined magnesia, 119
**Calcium**, 105
analytical reactions of, 114
and barium, separation from magnesium, 124
bisulphite, 306
carbonate of, 107
prepared, 110
chloride of, 105
chromate of, 115
citrate of, 326
derivation of word, 32
-flame, 115
fluoride of, 105
fluoride of, in bones, 111
gummate of, 114
hydrate of, 107
hypochlorite of, 113
hypophosphite of, 343
hyposulphite of, 344
in presence of barium and magnesium, detection of, 124
oxalate of, 115, 317
oxide of, 106
phosphate of, 105, 110
polysulphide of, 303
**Calcium—**
quantitative estimation of, 574
silicate of, 105, 353
sulphate of, 105, 114, 304
sulphite of, 306
superphosphate of, 327
tartrate of, 322
Calc-spar, 105
Calculi, urinary, 504
urinary, examination of, 504
Calomel, 195, 202
test for corrosive sublimate in, 202
tests for constituents of, 202
*Calotropis*, 400
*Calumbæ radix*, 394
*Calx*, 106
impurities in, 617
*Calx chlorata*, 113
*chlorata*, impurities in, 617
*chlorinata*, 113
*Cambogia*, 472
impurities in, 617
Camphor laurel, 467
mixture, 467
oil, 467
water, 467
*Camphora*, 467
impurities in, 617
*officinarum*, 467
Camphoretic acid, 467
Camphoric acid, 467
peroxide, 461
Camphors, 467
Camwood, 476
Canada balsam, 471
Canadian turpentine, 466
Candle-flame, composition of, 23
*Cannellæ albæ cortex*, 400
Cane-sugar, 410
Cannabene, 469
hydride of, 469
Cannabin, 469
*Cannabis indica*, 469
*Cantharides*, 468
Cantharidic acid, 468
Cantharidin, 468
*Cantharis*, 468
Caoutchouc, 474
Capacity unit, 521
Capillary, 513
Caproate of glyceryl, 458
Caproic acid, 458, 460

Caproyl, 433
Caprylate of glyceryl, 458
Caprylic acid, 458, 460
Capsaicin, 395
*Capsici fructus*, 395, 469
Capsicia, 395
Capsicin, 471
Capsicine, 395
Capsicum, 395
fruit, resin of, 395
oil of, 395
Caramel, 412
Caraway oil, 464
Carbamate of ammonium, 92
Carbazotic acid, 445
Carbolic acid, 444, 515, 517
acid, antidote to, 444
*Carbo animalis*, 111
*animalis purificatus*, 111
*animalis purificatus*, impurities in, 617
*ligni*, 112
*ligni*, impurities in, 617
**Carbon**, 30
bisulphide of, 433
combustion of, 30
derivation of word, 31
quantitative estimation of, in organic compounds, 596 *et seq.*
Carbonate of ammonium, 92
ammonium, solution of, 93
barium, 104
bismuth, 253
cadmium, 249
calcium, 105, 107
prepared, 110
iron, 143, 562
iron, saccharated, 143, 562
lead, 210
lithium, 227
magnesium, 117
potassium, 61, 547
potassium, acid, 71, 547
chemically pure, 550
sodium, 81, 87, 313, 547
sodium, acid, 83, 547
chemically pure, 550
sodium, manufacture of, 87, 313
strontium, 228
zinc, 130, 132
**Carbonates**, 312
acidulous radical in, 312
**Carbonates**—
analytical reactions of, 314
gravimetric estimation of, 591
volumetric estimation of alkaline, 547
Carbonic acid, 30, 312
acid gas, generation of, 71
acid gas, solubility of, in water, 85
anhydride, 312
oxide, 317, 338
Carbonization, 101
Cardamom oil, 464
Cardamomum, 464
greater, 464
lesser, 464
Carmine, 337
Carminic acid, 337
Carnallite, 61
Carolina jasmine, 396
Carrageen moss, 405
Carrotin, 475
Carthamin, 476
*Carthamus tinctorius*, 476
*Carum*, 464
*ajowan*, 463
Carvene, 464
Carvol, 464
Caryophyllin, 464
Cascarilla oil, 464
*Cascarilla cortex*, 464
Casein, 451
vegetable, 453
*Cassia marilandica*, 415
Cassia oil, 464
*Cassiæ pulpa*, 410
*Castilloa elastica*, 474
Cast-iron, 141, 517
Castor, 469
*Castor fiber*, 469
Castor oil, 459
*Castoreum*, 469
Castorin, 469
Catechin, 357
Catechuic acid, 357
Catechu, 357
*Catechu pallidum*, 357
*pallidum*, impurities in, 617
Cathartic acid, 415
Cathartogenic acid, 415
*Caulophyllum tralictroides*, 400
Caustic, 218
lime, 106
lunar, 218

Caustic—
potash, 62
soda, 81
Cedra oil, 463
Celestine, 228
Cellulin, 407
Cellulose, 407
Celsius's thermometer, 514
Cements, 353
Centiare, 522
Centigrade thermometer, 514
*Cephaelis ipecacuanha*, 396
*Cera alba*, 458
*alba*, impurities in, 617
*flava*, 458
*flava*, impurities in, 617
Cerasin, 405
*Cerasus virginiana*, 415
*Ceratum plumbi subacetatis*, vide *Unguentum*.
*zinci carbonatis*, 133
Cerebrin, 451
Ceresine, 458
*Cerii oxalas*, 229
*oxalas*, impurities in, 617
Cerite, 229
Cerium, 229
derivation of word, 33
oxalate of, 229
Ceroleine, 458
Cerotic acid, 458
*Cervisiæ fermentum*, 422
Cetaceum, 458
impurities in, 617
Cetene, 458
*Cetraria*, 337
Cetraric acid, 337
Cetyl, hydrate of, 458
palmitate of, 458
*Cevadilla*, 399
Cevadilline, 399
Cevadine, 399
Ceylon moss, 405
Chalcedony, 352
Chalk, 110
precipitated, 107
prepared, 110
-stones, 504
Chalybeate water, 140
Chameleon mineral, 231
Chamomile oil, 463
Char, 101
Charcoal, 29
animal, 111
Charcoal—
animal, decolorizing power of, 112
wood, 112
Charles's law, 52
*Chartæ sinapis*, 447
Chartreuse, 423
*Chavicæ officinarum*, 397
Chavicic acid, 398
Cheese, 451
Chemical action, definition of, 36, 40
action by symbols, illustration of, 41, 46
affinity, 38
combination, 38
combination by weight, laws of, 47, 58 *et seq.*
combination by volume, laws of, 52 *et seq.*
combination different from mechanical, 36, 38
combination, laws of, 47, 197, 286, 510
compound, definition of, 57
compound, 30
diagram, 57, 62
equation, 57, 62
force, 37, 56
force, conditions for the manifestation of, 40
force, its relations to heat and electricity, 539
formula, definition of, 57
formulæ, 41, 42
notation, 40, 41, 42
philosophy, principles of, 36 *et seq.*
preparations of the Pharmacopœias, 507
symbol, definition of, 57
symbols, 31, 40, 41, 42
toxicology, 483
Chemicals, 14
list of, xiii.
Chemism, 38
Chemist and Druggist, 14
Chemistry, art of, 13
and physics, differences between, 46
definition of, 56
derivation of the word, 13
inorganic, 378
object of, 14, 48

Chemistry—
organic, 378
science of, 13
Chemists, analytical, 14
manufacturing, 14
pharmaceutical, 14
*Chenopodium*, 467
Cherry-laurel water, 414
Cherry sugar, 410
Cherry-tree gum, 405
Cherry, wild black, 415
Chestnut-brown, 477
Chian turpentine, 466
Chili saltpetre, 285
*Chimaphila*, 415
*Chimaphila umbellata*, 415
Chinese red, 476
yellow, 475
Chinoidin, 603
*Chirata*, 350
Chiratin, 350
Chiratogenin, 350
*Chloral*, 438
Chloral, alcoholates of, 440
butyl, 440
croton, 431
hydrate, 438
*Chloral hydras*, 440
*hydras*, impurities in, 617
Chlorate of potassium, 293
potassium, preparation of oxygen from, 16
Chlorates, 293
analytical reactions of, 295
Chloric acid, 292
Chloride of ammonium, 90
antimony, 179
arsenicum, 167
barium, 102
boron, 333
calcium, 105
calcium, removal of iron from, 106
chromium, 239
gold, 246
iron, 147
lead, 213
lime, 113
magnesium, 117
manganese, 230
mercuric ammonium, 206
mercurous ammonium, 206
mercury, 200
platinum, 247
Chloride of—
platinum and ammonium, 98, 248, 572
platinum and lithium, 228
platinum and potassium, 78, 248
silicon, 354
silver, 217
sulphur, 305
tin, 242
solution of, 242
zinc, 131
Chlorides, 266
estimation of, 585
separation of, from bromides and iodides, 276
tests for, 269
Chlorinated lime, 113
lime, volumetric estimation of, 564
soda, solution of, 86
soda, volumetric estimation of, 564
**Chlorine**, 27, 268, 585
acids, 295
as a disinfectant, 29
bleaching by, 28
collection of, 27
derivation of word, 31
its analogy to bromine and iodine, 273, 276
liquid, 268
percentage in hydrochloric acid, 628
preparation of, 28
properties of, 27
relative weight of, 28
solubility in water, 28
the active agent in bleaching-powder, 113
volumetric estimation of, 565
water, 28, 268
Chlorochromic anhydride, 239
Chloroform, 436, 515
impurities in, 617
water, 438
Chlorophyll, 477, 507
Chocolate, 458
Cholalic acid, 457
Cholate of sodium, 457
Cholesterin, 451, 506
Chondrin, 453
*Chondrus crispus*, 405
Chromate of barium, 104

Chromate of—
calcium, 115
lead, 214
mercury, 239
potassium and ammonium, 104
potassium, standard solution of red, 561
silver, 219
Chromates, 238
analytical reactions of, 240
of potassium, 104, 124, 237
Chrome-alum, 238
-ironstone, 237
-red, 214
-yellow, 214
Chromic acid, 238
anhydride, 238
hydrate, 238
salts, 238
**Chromium**, 237
analytical reactions of, 240
chloride of, 238
derivation of word, 33
separation of, from aluminium and iron, 240
sulphate of, 238
Chromous salts, 240
Chromule, 477
Chrysammic acid, 446
Chrysarobine, 337
Chrysophanic acid, 337
*Cicuta virosa*, 464
Cider, 423
*Cimicifuga racemosa*, 400
Cimicifugin, 400
*Cinchona Calisaya*, 384
*Cinchonæ flavæ cortex*, 384
*officinalis*, 384
*paillidæ cortex*, 384
*rubræ cortex*, 384
*succirubra*, 384
Cinchonia, 388
*Cinchoniæ sulphas*, 388
Cinchonicia, 388
Cinchonicine, 388
Cinchonidia, 388
Cinchonidine, 388
Cinchonine, 388
Cinnabar, 194
Cinnamein, 473
Cinnamene, 473
Cinnamic acid, 473
alcohol, 473
Cinnamic—
aldehyd, 464
Cinnamol, 473
*Cinnamomi cortex*, 464
Cinnamon oil, 464
Cinnamyl, hydride of, 473
alcohol, 473
cinnamate of, 473
**Citrate** of ammonium, 94
calcium, 324
iron, 152
iron and ammonium, 153
iron and strychnia, 152
iron and quinine, 152, 606
lithium, 227
magnesium, 120
nicotia, 397
potassium, 72
volumetric estimates of, 547
quinine, 386
silver, 326
Citrates, 72, 324
analytical reactions of, 326
volumetric estimation of, 547, 551
Citric acid, 324
action of heat on, 326
saturating power of, 325
Citronella oil, 464
Citronellol, 464
Citro-tartrate of sodium, 86
*Citrus*, 464
*Citrus* bergamia, 324
Classification of elements, 101, 122, 125, 259
Clausius's theory, 24
Claviceps purpurea, 470
Clay, 137
ironstone, 140
Cloves, oil of, 464
Club-moss, 459
Coal, 241
products of, 433
Coal-gas, 433
for balloons, 23
-tar colors, 478
**Cobalt**, 234
analytical reactions of, 234
arsenide of, 234
blue, 477
derivation of word, 33
-glance, 234
hydrate of, 234

**Cobalt—**
separation of, from nickel, 236
sulphate of, 234
sulphide of, 234
Cobaltic ultramarine, 477
Cobalticyanide of potassium, 235
of nickel, 235
Cobalticyanides, 235
Coca, 386
Cocaina, 386
Cocaine, 386
*Cocculus indicus*, 420
*Coccus*, 337
*Ilicis*, 182
Cochineal, 337
Cocoa, 458
Cocoa-butter, 458
Cocoa-nibs, 458
Cocoa-nut, 458
Cocoa-nut oil, 458
*Cocos nucifera*, 458
Codamine, 382
Codeia, 382
Cod-liver oil, 459
Cohesion, 55
Coinage, copper, 191
gold, 245
silver, 216
Coke, 30
*Colchici cormus*, 395
*semina*, 395
Colchicia, 395
Colchicin, 395
Colchicine, 395
Colcothar, 476
Collection of gases, 16, 17
Collin, 611
Collodion, 408
*Collodium*, 408
*flexile*, 408
Colloid bodies, 596
*Colocynthidis pulpa*, 416
Colocynthin, 416
Colopholic acid, 469
Colophonic acid, 469
hydrate, 469
Colophonine, 469
Colophony, 469
Coloring-matters, 475
Combination, chemical, 44, 46
by volume, 52
Combining proportions, 48, 197, 286, 511, 568
Combustible, 22
Combustion, 22
analysis for carbon and hydrogen, 596 *et seq.*
for nitrogen, 600
definition of, 57
spontaneous, 155
supporters of, 22
Composition of atmosphere, 26
bismuth salts, 253
oils and fats, 456
Compound, chemical, 37
different from mechanical, 36
definition of, 57
Compounds, 13, 36
of the elements, 60
Conchinine, 388
Condensation, 126
Condenser, 127
Condensing-tub, 127
worm, 127
Condy's disinfecting fluids, 77 231
Confections, 507
Conhydrine, 396
Conia, conine, or conicine, 395
*Conii folia*, 395
*fructus*, 395
Conium maculatum, 395
Conquinine, 388
Constant proportions, law of, 47
Constitution of alkaloids, 379
of matter, 41
of salts, 52, 123, 263, 282, 300, 377
of visible matter, 30, 42
Construction of formulæ, 41, 45, 62, 63, 406
Convolvulin, 419
Convolvulinol, 419
*Convolvulus scammonia*, 421
Conylia, 395
*Copaiba*, 471
impurities in, 471, 617
Copaiva, 471
oil, 471
Copaivic acid, 471
Copal, 469
**Copper**, 190
acetate of, 192
ammonio-sulphate of, 177, 193, 206
analytical reactions of, 192, 485

**Copper—**
antidotes to, 193
arseniate of, 177
arsenical, 173
arsenite of, 177
black oxide of, 191
blue, 477
coinage, 191
derivation of word, 32
detection of arsenicum in, 177
foil, 191
hydrate of, 193
iodide of, 276
in organic mixtures, detection of, 484
melting-point of, 517
metallic, 191
oxide of, 191
oxyacetate of, 192
pyrites, 190
quantitative estimation of, 580
quantivalence of, 191
subacetate of, 192
sulphate of, 192
sulphate of, anhydrous, 192
sulphide of, 192
Copperas, blue, 143
green, 143
*Coptis trifolia*, 394
Coriander oil, 464
*Coriandri fructus*, 464
*Coriandrum*, 464
Cork-borers, 16
Corpse-fat, 453
Correction of the volume of a gas for pressure, 536
for temperature, 536
Corrosive sublimate, 196, 200
test for, in calomel, 202
Cortex pruni serotinæ, 415
Corydalia, 396
Corydalina, 396
Cotarnine, 382
Coto bark, 416
Cotoin, 416
Cotton-wool, 407
Cow-bane, 464
Cowhage, 241
Cow's milk, 452
Cranesbill, spotted, 357
Cream, 452
Cream of tartar, 319
Creasol, 444
Creasote, 444
*Creasotum*, 444
impurities in, 617
Cresol, 444
Cresotic acid, 445
Cresylic acid, 444
*Creta*, 110
*præparata*, 110
*Crinum asiaticum*, 422
Crocin, 475
Crocus (mineral), 152
(vegetable), 475
*Crocus sativus*, 475
Croton chloral, 441
Croton oil, 459
Crucibles, 69
Crude antimony, 179
potashes, 61
Crum's test for manganese, 233
Cryohydrates, 85
Cryolite, 375
Cryptophanic acid, 494
Cryptopia, 382
Crystallization, water of, 84
Crystalloid bodies, 611
*Cubeba*, 397
Cubebene, 464
Cubebin, 464
Cubeb pepper, 397
Cubebs, oil of, 464
Cubic inches in one gallon, 537
Cubic nitre, 285
Cuca (*see* Coca).
Cumin, 464
Cuminic acid, 465
Cuminol, 464
*Cuminum*, 464
*cyminum*, 464
Cummin, 464
Cupel, 584
Cupellation, estimation of silver by, 584
Cupr-diammon-diammonium, sulphate of, 206
*Cupri subacetas*, 192
*sulphas*, 192
*sulphas*, impurities in, 617
Cepric arsenite, 177, 193
compounds, 192
ferrocyanide, 193
hydrate, 193
oxide, 191

Cupric—
sulphate, 192
sulphide, 192
Cuprous iodide, 191, 276
oxide, 192, 409
*Cuprum*, 191
*ammoniatum*, 193
Curaçoa, 423
Curarine, 391
*Curcuma longa*, 475
Curcumin, 475
Curd-soap, 457
Curds, 411, 451
and whey, 411, 451
Currant, sugar in, 410
*Cusparie cortex*, 400
Cusparin, 400
*Cusso*, 470
Cutch, 357
Cyanates, 337
Cyanic acid, 337
Cyanide of allyl, 448
mercury, 280
nickel, 236
potassium, 279
silver, 219
Cyanides, 279
analytical reactions of metallic, 282
antidote to, 283
double, 279
quantitative estimation of, 556
**Cyanogen**, 279, 286
Cyanurets, *vide* Cyanides.
Cyder, *see* Cider.
*Cydonia vulgaris*, 406
Cymene, 464
Cymol, 464
*Cypripedium pubescens*, 400
Cystin, 499

Dahlia, 404
Dalton's atomic theory, 50
law, 48, 49
Danglish's bread, 411
*Daphne laureola*, 470
*mezereum*, 416, 470
Daphnetin, 416
Daphnin, 416
*Datura alba*, 393
Datura or daturine, 393
Davy's safety-lamp, 23
Davyum, 635
Deadly nightshade, 393
Decantation, 109
Decimal coinage, 520
weights, 520, 523
Decoctions, 507
Decolorizing power of animal charcoal, 112
Decrepitation, 370
Definition of chemical action, 36
a chemical compound, 57
a chemical equation or diagram, 57
a chemical symbol, 57
a gas, 58
a liquid, 57
a mixture, 57
an atom, 57
an element, 57
a solid, 57
atomic weights, 58
chemical force, 56
chemistry, 56
combustion, 57
law of diffusion, 57
molecular weights, 58
quantivalence of atoms, 58
Deflagrating flux, 375
Deflagration, 74
Deliquescence, 87
Delphine, 396
Density, 521
of vapors, 536
Dentifrices, action of, 241
Deodorizers, 29
Deodorizing liquid, 131
Deposits, urinary, 497
Derivation of names of elements, 31 *et seq.*
Derivatives of ammonium, 206
Desiccation, 569, 591
Destructive distillation, 127
Detonation, 74
De Valangin's solution, 168
De Vrij's process for estimating the value of cinchona bark and the purity of commercial quinine, 601
Dextrine, 404
maltose, 412
Dextroracemic acid, 320
Dextrose, 412
Dextrotartaric acid, 320
Dhak tree, 357
Dhatura, 393

Diabetic urine, 495
Diagram, chemical definition of, 57
Diagrams, chemical, 46, 47, 63, 65, 70
Dialysate, 611
Dialysis, 610
Dialytic iron, 612
Dialyzed iron, 612
Diamines, 379
Diamond, 29
Diastase, 405
Dibasic acids, 265
Dibasylous radicals, 265
*Dicentra formosa*, 396
Didymium, 635
Diethylamine, 379
Diethylia, 379
Diffusate, 611
Diffusion, 24
  law of, definition of, 57
Digitalein, 417
Digitalin, 416
*Digitalinum*, 416
Digitaliretin, 416
*Digitalis folia*, 417
Digitonin, 417
Digitoxin, 417
Dill oil, 463
Dinitrobenzol, 444
Dinitrocellulin, 408
Diospyros embryopteris, 357
*Dipterocarpus lævis*, 471
  *turbinatus*, 471
Disinfectant, chlorine as a, 29
Disinfectants, 29
Disinfecting fluid, Burnett's, 131
    carbolic acid, 444
    Condy's 77, 231
  powder, 113
Dissociation, 539
Distillation, 126
  destructive, 127
  dry, 127
  fractional, 423
Distilled vinegar, 298
Dita, 396
Ditain, 396
Ditamine, 396
Dithionic acid, 345
Dock, 337
Dolomite, 116
Donovan's solution, 198
*Dorema ammoniacum*, 472
Double chloride of aluminium and sodium, 137
  cyanides, 279
  salts, 78
Dover's powder, 396
Drachm, 526
Draconyl, 474
Dragon's blood, 469
Dried alum, 138
Drops, 526
Dry distillation, 127
Drying apparatus, 568, 570, 591
  oils, 459
  precipitates, 567, 569, 573, 591
*Dryobalanops aromatica*, 467
Duboisia, 396
*Dulcamara*, 398
Dulcamarin, 398
Dulong and Petit's law, 539
Dyads, 122
Dyer's saffron, 476
Dyeing, 139, 291
  by mordants, 139
Dynamic electricity, production of, 131
Dynamicity, 56

Earth, bone, 110
Earth-nut oil, 459
Earthenware, 353
Earth, alkaline, 126
Eau de Cologne, 463
Ebonite, 474
Ebullition, 281
*Ecballii fructus*, 417
Ecboline, 470
*Echites scholaris*, 396
"Effervescing citrate of magnesia," 120
  citro-tartrate of sodium, 86
  potash-water, 72, 571
  soda water, 85
Efflorescence, 87
Egg, oil of, 450
  white of, 449
  yolk of, 450
Elæometer, 533
Elæoptens, 461
Elaterin, 417
*Elaterium*, 417
  impurities in, 617
Elder-flower oil, 466
Elecampane, 404, 465

Electricity, production of, dynamic, 131
related to chemical action, 539
Elementary particles, 37
Element, definition of, 57
Elements, 13, 14, 30
and their compounds, 60
classification of, 101, 122, 125, 259
classification of, according to analogy, 101
classification of, according to quantivalence, 121
etymology of names of, 30
of medical or pharmaceutical interest, 14
metallic, 15
non-metallic, 15
of pharmaceutical interest, 14
symbols of, 31, 46
symbols, atomic values and weights of the, 635
symbols of, and derivation names of, 31 *et seq.*
Elemi, 471
Elm, common, 406
Elm mucilage, 406
Elm, slippery, 406
Elm tannin, 357
Emetia, 396
Emetine, 396
Empirical formulæ, 424
*Emplastra*, 469
*Emplastrum cerati saponis*, 298
*plumbi*, 213
*plumbi iodidi*, 212
Emulsin, 414
Emulsion, 473
Enemas, 507
English red, 476
blue, 477
Epsom salt, 116
Equation, chemical, definition of, 57
Equations, 57
Equisetic acid, 325
Equivalence, 56
Equivalents, 635
Ergot, 470
*Ergota*, 470
Ergotin, 470
Ergotine, 470
Ergotinine, 470
Ericolin, 415
Erlangen blue, 427
Erucic acid, 459
Erythroretine, 337
Erythroxylon coca, 395
Esculin (*see* Æsculin).
Eseria, 397
Essence of aniseed, 463
apple, 441
greengage, 441
melon, 442
mirbane, 443
mulberry, 442
peppermint, 441
pineapple, 441
quince, 441
Essences, 441
*Essentia anisi*, 462
*menthæ piperitæ*, 462
Essential oils (*vide* Oils).
Estimation of weight, 518
Etching, 341
Ethal, 458
Ether, 427
nitrous, 430
Ethereal salts, 433
oil, 447
Etherol, 447
Ethers, 427, 433
Ethiops mineral, 207
**Ethyl**, 424, 432
acetate, 431
bromide, 432
butyrate, 441
hydrate, 422
hydride, 432
iodide, 432
nitrite, 430
œnanthylate of, 441
oxide, 427
pelargonate of, 441
sebacate, 442
suberate, 441
sulphate, 424
zinc, 432
Ethylamine, 379
Ethylene, 447
Ethylia, 379
Ethylic alcohol, 422
Etymology of names of elements, 31
Eucalyptol, 465
*Eucalyptus globulus*, 465
Euchlorine, 296

Eudiometry, 481
Eugenic acid, 464
Eugenin, 464
Euodic aldehyd, 466
Euonymin, 400
*Euonymus Europæus*, 400
Euphorbium, 472
Euphorbon, 472
Euxanthate of magnesium, 475
Evaporation, 70, 100, 569, 570
*in vacuo*, 570
Everitt's yellow salt, 281
Examinations of the pharmaceutical Society of Great Britain, 14 (*vide* Prefatory matter).
Expansion on diluting solution of ammonia, 91
Explosions of gases, 22
Extracts, 507
*Extractum cannabis*, 469
*filicis liquidum*, 459
*glycyrrhizæ*, 418
*liquidum*, 418
*Saturni*, 211

Face-rouge, 337, 476
Fahrenheit's thermometer, 514
*Farina tritici*, 401
Fat-acids, 460
Fats and oils, composition of, 456
Fats, etc., to determine the melting point of, 516
analysis of, 613
Fatty bodies, 456
Feces, 493
*Fel bovinum purificatum*, 457
*purificatum*, impurities in, 617
Felspar, 375
Fennel oil, 465
Fermentation, 410
*Fermentum*, 422
*Fer réduit*, 158
Ferrate of potassium, 141
*Ferri acetas tinctura*, 149
*arsenias*, 144
*arsenias*, impurities in, 618
*carbonas*, 143
*carbonas saccharata*, 143
*carbonas saccharata*, impurities in, 618
*chloridi, liquor*, 148
*chloridum*, 148
*citras*, 152
*et ammoniæ citras*, 153
*Ferri*—
*et ammoniæ citras*, impurities in, 618
*et ammoniæ citras*, quantitative estimation of iron in, 578
*et ammoniæ sulphas*, 138
*et ammoniæ tartras*, 155
*et potassæ tartras*, 155, 323
*et quiniæ citras*, 153
*et quiniæ citras*, impurities in, 618
*et strychniæ citras*, 154
*ferrocyanidum*, 340
*hypophosphis*, 343
*iodidum*, 31, 146
*lactas*, 346
*oxalas*, 316
*oxidum magneticum*, 156
*oxidum magneticum*, impurities in, 618
*perchloridi, liquor*, 148
*pernitratis, liquor*, 157
*peroxidum humidum*, 150
*peroxidum hydratum*, 150
*peroxidum hydratum*, impurities in, 618
*persulphatis, liquor*, 153
*phosphas*, 144
*phosphas*, impurities in, 618
*potassio-tartras*, 155
*pulvis*, 158
*pyrophosphas*, 159
*subcarbonas*, 144
*subsulphatis, liquor*, 149
*sulphas*, 142
*sulphas*, impurities in, 618
*sulphas exsiccata*, 142
*sulphas granulata*, 142
*sulphas granulata*, impurities in, 618
*sulphuretum*, 145
**Ferric** acetate, 149
chloride, 146
citrate, 152
hydrate, 150
hypophosphite, 342
iodate, 296
nitrate, 157
oxide, 151
oxide, from phosphates and oxalates, separation of, 373
oxyiodate, 296
oxysulphate, 143
peroxyhydrate, 150

**Ferric—**
pyrophosphate, 157
salts, 146
salts, analytical reactions of, 160
sulphate, 149
sulphocyanate, 161
tartrate, 155
Ferridcyanide of potassium, 160, 279, 340
Ferridcyanides, 340
Ferrocyanogen, 161, 340
Ferrocyanide of potassium, 160, 279, 339
of zinc, 135
Ferrocyanides, 339
Ferrocyanogen, 161, 339
Ferro-ferric hydrate, 156
oxide, 156
**Ferrous** arseniate, 144, 561
bromide, 146
carbonate, 143, 561
chloride, 147
hydrate, 160
iodide, 146
phosphate, 144
salts, 142
salts, analytical reactions of, 159
sulphate, 142
sulphide, 145, 160
*Ferrum*, 32, 140
*redactum*, 157
*tartaratum*, 155, 323
*tartaratum*, estimation of iron in, 578
*tartaratum*, impurities in, 618
Ferulaic acid, 472
Fibrin, 450
vegetable, 453
*Ficus*, 410
*elastica*, 474
Fig, 410
Filicic acid, 459
*Filix mas*, 459
Filter, to dry, 568
Filtering-paper, 108, 568
Filters, 108, 568
Filtrate, 124
Fine gold, 246
Fire-clay, 201
Fire-damp, 433
Fixed and volatile oils, difference between, 459
Fixed—
oils, 456
Flame, structure of, 23
Flaxseed, 459
Fleitmann's test for arsenicum, 174
Flexible collodion, 408
Flint, 352
*Flores zinci*, 134
Flour, 401
Flowers of sulphur, 302
Fluid magnesia, 118
Fluoride of boron, 333
calcium, 105
calcium, in bones, 111, 342
silicon, 342
Fluorides, 341
Fluorine, 342
derivation of word, 33
Fluor-spar, 341
*Fœniculi fructus*, 465
Foil, copper, 191
Food, analysis of, 613
elements of, 453
how disposed of in the bodies of animals, 493
Force, chemical, 37
Forge-scales, 156
Formates, 338
Formic acid, 337, 460
*Formica rufa*, 337
Formula, chemical, definition of, 57
official, 28
officinal, 28
Formulæ, 41, 45, 601
construction of, 55, 62
empirical, 424, 601
graphic, 136
rational, 424, 601
typical, 379
Fousel oil, 441
Fowler's solution, 167
Foxglove, 416
Fractional distillation, 423
Frankincense, Arabian, 473
common, 471
Frankland's graphic formulæ, 136
*Fraxinus ornus*, 413
Free acids estimated, 552
Freezing-mixture, 305
French chalk, 478
turpentine, 466
Fruit essences, 441

Fuchsine, 478
Fume-cupboard, 96
Fuming sulphuric acid, 310
Funnel-tubes, 23, 97
"Fur" in water vessels, 315
Furniture of a laboratory, xii.
Fusel oil, 441
Fusibility of metals, table of the, 517
Fusible white precipitate, 206
Fusing-points of fats, 516
Fustic, 475

Gab tree, 357
Galactometer, 533
*Galbanum*, 472
Galena, 209
  argentiferous, 216
Galenical preparations of the British Pharmacopœia, 507
Galipot, 469
*Galla*, 355
Gallic acid, 357
Gallium, 635
Gall of the ox, 457
Gallon, 520
Galls, Aleppo, 355
  English, 355
Gall-stones, 506
Galvanic test for mercury, 204
Galvanized iron, 130
Gambier, 357
Gamboge, 472, 475
Gambogic acid, 472
Garancin, 476
*Garcinia indica*, 459
  *indica*, oil, 459
  *morella*, 472
  *pictoria*, 472
  *purpurea*, 459
Garden thyme, 465
Garlic, essential oil of, 447
Gas, a, definition of, 57
  -analysis, 358, 481
  -burners, 16, 23
  for balloons, coal-, 23
  illuminating, 433
  -lamp, 18, 23
Gases and vapors, density of, 512, 536
  collection of, 16, 19
  correction of the volume of, 536
  for pressure, 536
Gases, correction—
    for temperature, 536
  diffusion of, 24
  law of solubility of, in liquids, 85
  relation of, to liquids and solids, 43, 44
  specific gravity of, 536
Gastric juice, 454
    artificial, 454
*Gaultheria procumbens*, 442
Gaultheric acid, 442
Gay Lussac's law, 52
Gelatine, 453
  sugar, 457
  vegetable, 405
Gelatinous substances, 453
Gelseminia, 396
Gelseminic acid, 396
Gelsemium, 396
Gentian-bitter, 418
*Gentiana lutea*, 418
*Gentianæ radix*, 418
Gentianic acid, 418
*Gentiogenin*, 418
Gentiopicoris, 418
Gentisic acid, 418
Gentisin, 418
Geraniol, 465
*Geranium maculatum*, 357
German silver, 235
Gin, 423
Gingelly oil, 460
Ginger oil, 467
  -grass oil, 465
Girdwood and Rogers's method for detecting strychnia, 489
Glacial acetic acid, 299, 517
  phosphoric acid, 330
Glass, 353
  liquor, 353
  rods, 109
  soluble, 353
  tubes, to bend, 17
    to cut, 17
    to draw out, 109
Globulin, 451
Glucinum, 635
Glucose, 410
Glucosides, 413
Glue, 454
Gluten and glutin, 401
Glyceric alcohol, 448
Glycerine, 213, 448

Glycerines, 449
*Glycerina*, 449
impurities in, 618
*Glyceritum acidi carbolici*, 449
*acidi gallici*, 449
*acidi tannici*, 358, 449
*picis liquidæ*, 449
*sodii boratis*, 449
Glyceryl, 448, 456
caproate, 458
caprylate, 458
hydrate, 448, 456
laurate, 458
myristate, 458
oleate, 456
palmitate, 458
ricinoleate, 458
rutate, 458
tristearate, 456
Glycholates, 457
Glycocine, 457
Glycocoll, 457
Glycol, 447
Glycols, 447
Glycyl, 448
Glycyrretin, 418
*Glycyrrhizæ radix*, 418
Glycyrrhizin, 418
Gnoscopine, 382
Goa powder, 337
**Gold**, 244
analytical reactions of, 246
coin, 245
derivation of word, 34
earth, 244
fine, 246
jewellers', 245
leaf, 245
mosaic, 244
ochre, 475
perchloride of, 245
sulphide of, 246
yellow, 475
Golden seal, 394
Goldthread, 394
*Gossypium*, 407
Gothite, 151
Goulard's cerate, 211
extract, 211
water, 211
*Gracellaria*, 405
Graham's dialytic process, 610
law of diffusion, 24
Grain, 519
Grains, 519
of paradise, 465
Gramme, 522
relation of, to grains, 524, 525
*Granati fructus cortex*, 357
*radicis cortex*, 357
Granite, 137
Granulated phosphorus, 328
tin, 241
zinc, 20
Grape-sugar, 410
Grape, sugar in, 410
Grapes, dried, 410
Graphic formulæ, 137
Graphite, 29
Grass oil, 464, 467
Gravel, 496
Gravimetric analysis, 510, 567
Gravitation, 518
Gravity, 518
Gray powder, 195
Green copperas, 143
chloride of iron, 147
iodide of iron, 146
iodide of mercury, 196
pigments, 477
sulphate of iron, 142
vitriol, 143
Greengage essence, 441
Griffith's mixture, 144
Groundnut oil, 459
Group tests, 224
*Guaiaci lignum*, 418
*resina*, 418
Guaiacin, 418
Guaiacol, 444
Guaiaconic acid, 418
Guaiacum, resin of, 418
Guaiaretic acid, 418
Guaiaretin, 418
Guaiaretinic acid, 418
Guano, 359
Guarana, 398
*Guilandina bonducella*, 400
Guinea grains, 465
*Gulancha*, 400
Gum, 405
-acacia, 114
-arabic, 114
British, 404
cherry-tree, 405
-resins, 472
tragacanth, 114, 405
Gummate of calcium, 114, 405

Gummate of—
lead, 114
Gummic acid, 405
Gun-cotton, 408
Gun metal, 241
Gunpowder, 290
Gurjun balsam, 471
Gutta percha, 474
Guttæ, 526
*Gynocardia odorata*, 400
Gypsum, 105

Hæmatein, 477
Hæmatite, brown, 140
red, 140
*Hæmatoxyli lignum*, 357, 476
Hæmatoxylin, 357, 476
Half-sovereign, weight of the, 245
Haloid salts, 287
Hambro' blue, 477
Hardness of water, 315
Hard soap, 457
Heat, latent, 84
related to chemical action, 539
source of, 21
Heavy carbonate of magnesium, 118
magnesia, 119
spar, 102
white, 102, 478
Heberden's ink, 161
Hectare, 522
Helenin, 465
Hellebore, black, 419
green, 419
white, 419
American, 397
Helleborein, 419
Helleborin, 419
*Helleborus niger*, 419
*viridis*, 419
*Hemidesmi radix*, 338
Hemidesmic acid, 338
Hemlock, 395.
Hempseed calculi, 506
Henbane, 396
Henry and Dalton's law, 85
Herapathite, 387
Hesperidene, 463
*Hevea siphonia Braziliensis*, 474
Heterologous series, 434
Hexyl, 433
*Hibiscus esculentus*, 406
Hippuric acid, 338, 497, 500
Hips, 412
Hoffner's blue, 477
Homologous series, 434, 460
Honey, 412
dew, 412
Hop, 472
essential oil of, 472
*Hordeum decorticatum*, 401
starch of (fig.), 403
Horehound, *vide* Marrubium.
Horsemint, 466
Horseradish oil, 434, 463
Hubbuck's oxide of zinc, 134
*Humulus lupulus*, 472
*Hydrargyri chloridum corrosivum*, 200
*chloridum mite*, 202
*cyanidum*, 280
*iodidum rubrum*, 198
*iodidum rubrum*, impurities in, 618
*iodidum viride*, 196
*iodidum viride*, impurities in, 618
*nitrati acidus, liquor*, 199
*nitratis, liquor*, 199
*oxidum flavum*, 203
*oxidum flavum*, impurities in, 618
*oxidum rubrum*, 203
*oxidum rubrum*, impurities in, 618
*perchloridum*, 200
*subchloridum*, 202
*subchloridum*, impurities in, 618
*sulphas*, 199
*sulphas flava*, 200
*sulphas*, impurities in, 618
*sulphuretum cum sulphure*, 207
*sulphuretum cum sulphure*, impurities in, 618
*sulphuretum rubrum*, 207
*unguentum*, 195
*Hydrargyrum*, 33, 194
impurities in, 618
*ammoniatum*, 205
*ammoniatum*, impurities in, 618
*cum creta*, 195
*cum creta*, impurities in, 618
Hydrastia, 394
*Hydrastis canadensis*, 394
Hydrated peroxide of iron, 150
substances, 84

Hydrate of aluminium, 139
ammonium, 91
benzoyl, 336
cadmium, 250
calcium, 107
cetyl, 458
chromium, 240
cobalt, 234
glyceryl, 458
manganese, 233
nickel, 236
potassium, 62
sodium, 81
zinc, 135
Hydrates, composition of, 62, 84
identified, 375
Hydraulic cement, 353
Hydric acetate, chloride, nitrate, sulphate, etc., *vide* the respective acids—acetic, hydrochloric, etc.
Hydride of antimony, 185
arsenicum, 174
benzoyl, 414, 472
cinnamyl, 473
copper, 344
ethyl, 432
methyl, 433
phosphorus, 343
silicon, 353
Hydrides, 122, 434
Hydriodic acid, 273
*Hydrium*, 20
Hydrobromic acid, 270
ether, 432
Hydrocarbons, 461
Hydrochlorate of morphia, 381
Hydrochloric acid, 29, 266
acid, analytical reactions of, 269
acid, antidote to, 269
acid, common, 266
dilute, 266
acid in organic mixtures, detection of, 486
acid in the liquid acid of different densities, Table showing the amount, 628
acid, volumetric estimation of, 554
Hydrocotarnine, 382
*Hydrocotyle asiatica*, 400
Hydrocyanic acid, 279
acid, analytical reactions of, 282, 486
Hydrocyanic—
acid, antidotes to, 283
acid, dilute, 280
acid, from bitter almond and cherry laurel, 414
acid in organic mixtures, detection of, 486
acid in the blood, 284
acid, Schönbein's test for, 284
acid, volumetric estimation of, 556
Hydroferridcyanic acid, 340
Hydroferrocyanic acid, 339
Hydrofluoric acid, 341
**Hydrogen**, 20
antimoniuretted, 185
arseniuretted, 174
benzoate, borate, etc., *vide* the respective acids, benzoic, boracic, etc.
combustion of, 21
derivation of word, 31
explosion of, 22
functions of, 122
in artificial light-producers, 22
lightness of, 23
peroxide of, 103
persulphide of, 303
phosphoretted, 343
preparation of, 20
properties of, 21
quantitative estimation of, in organic compounds, 599 *et seq.*
salts of, 263
siliciuretted, 353
sulphuretted, 96, 303
type, 377
used for balloons, 23
weight compared with air, 24
weight of 1 litre, 537
weight of 100 cubic inches, 537
*Hydrogenium*, 20
Hydroquinone, 415
Hydrometers, 533
Hydrosulphuric acid, 303
Hydrosulphyl, 303
Hydrous chloral, 439
Hydrous compounds, 84, 119
Hydroxyl, 303
Hyoscyamia, 396
*Hyoscyamia folia*, 396
*semen*, 396

Hyoscyamine, 396
Hyper-, meaning of, 146
Hypo-, meaning of, 343
Hypobromites, 271
Hypochloride of sulphur, 305
Hypochlorite of calcium, 113
sodium, 86
Hypochlorites, 292
Hypochlorous acid, 292
Hypophosphite of calcium, 343
of iron, 343
of potassium, 343
Hypophosphites, 342
Hypophosphoric acid, 351
Hypophosphorous acid, 342
Hyposulphite of calcium, 344
sodium, 344
sodium, standard solution of, 564
Hyposulphites, 344
Hyposulphurous acid, 344

-ic, meaning of, 76, 141
Icacin, 471
Iceland moss, 337
*Ichthyocolla*, 453
-ide, meaning of, 76
*Igasuria*, 391
*Ignatia*, 389
Ignition, 101
*Illicium anisatum*, 463
Illuminating agents, analysis, 613
Incense, 473
Inch, 526
Incineration, 101
of filters in quantitative analysis, 568, 573
Indelible ink, 218
Indian barberry, 394
gamboge, 472
hemp, 469
ink, 477
ipecacuanha, 396
mustard, 447
liquorice, 418
pennywort, 400
red, 476
rubber, 474
rubber, vulcanized, 474
yellow, 475
Indican, 290
Indiglucin, 291
Indigo, 290
Indigo—
sulphate of, 290
-blue, 290
-white, 291
wild, 400
Indigogen, 291
Indigotin, 291
Indium, 635
Infusible white precipitate, 206
Infusions, 507
Inhalation of chlorine, 28
conine, 395
hydrocyanic acid, 282
Inhalations, 507
*Injectio morphiæ hypodermica*, 381
Ink, black, 161, 356
Heberden's, 161
indelible, 218
Indian, 477
invisible, 235
marking, 218
printer's, 477
sympathetic, 235
Inorganic chemistry, 377
compounds, 377
Insecticide, 470
Introduction, 13
*Inula helenium*, 404, 465
Inulic anhydride, 465
Inulin, 404
Inulol, 451
Inverted sugar, 409
Iodal, 440
Iodate of potassium, 74, 296
Iodates, 296
Iodic acid, 296
Iodide of ammonium, 274
of arsenicum, 166
cadmium, 249
ethyl, 432
hydrogen, 274
iron, 31, 146
lead, 212
potassium, 73
silver, 219
sulphur, 274
**Iodides**, 273
analytical reactions of, 274
of mercury, 196, 205
quantitative estimation of, 586
separation of, from bromides and chlorides, 276
**Iodine**, 30, 273
chloride of, 273

**Iodine—**
derivation of word, 31
its analogy to chlorine and bromine, 278
solution of, 274
standard solutions of, 558
tincture of, 274
volumetric estimation of, 564
-water, 274
Iodoform, 425, 438
*Iodum*, 273
impurities in, 618
*Iodinium*, 273
*Ipecacuanha*, 396
*Ipomea orizabensis*, 419
*purga*, 419
*simulans*, 419
*turpethum*, 200
Iridin, 400
Iridium, 247, 248
*Iris florentina*, 465
*versicolor*, 400
Irish moss, 405
Irisin, 400
**Iron**, 140
acetate, 149
acetonitrate, 157
alum, 138
ammonio-citrate, 153
ammonio-tartrate, 155
analytical reactions, 159
arseniate, 144, 170
arseniate, volumetric estimation of, 562
black hydrate of, 156
black oxide of, 156
bromide, 146
carbonate, 143, 562
cast, 141
chlorides, 146
citrates, 152
compounds, nomenclature of, 142
derivation of word, 32
detection of, in presence of aluminium and zinc, 162
galvanized, 130
hydrated peroxide of, 150
hypophosphite of, 343
in official compounds, estimation of, 562, 578
iodate, 296
iodide, 30, 146
lactate, 346
**Iron—**
magnetic oxide of, 156
magnetic oxide of, estimation of iron in, 562
nitrate, 157
ore, magnetic, 140
ore, needle, 151
ore, spathic, 140
ore, specular, 140
oxide of, 156
oxyhydrates, 151
oxysulphate, 143
perchloride, 146
perhydrate, 150
pernitrate, 157
peroxide, 151
peroxyhydrate, 150
persulphate, 149
phosphate, 144
phosphate, volumetric estimation of, 562
from phosphates and oxalates, separation of peroxide of, 377
potassio-citrate, 152
potassio-tartrate, 155
protected from rust, 141
pyrites, 140
pyrophosphate, 159
quantitative estimation of, 562, 578
and quinine, citrate of, 153, 385
red oxide, 150
reduced, 157
rust, 141
saccharated carbonate, 143
saccharated carbonate, volumetric estimation of, 562
salts, nomenclature of, 141
scale compounds of, 152
separation of, from aluminium and chromium, 240
sodio-citrate, 152
sodio-tartrate, 152
-stone, clay, 140
subcarbonate, 144
subsulphate, 149
sulphate, 142
sulphide, 145
sulphocyanate, 161, 283
tartrate, 152
tersulphate, 149
wrought, 141

Isinglass, 453
Isomerides, 406
Isomerism, 406
Isomers, physical, 407
Isomorphism, the doctrine of, 54
Isomorphous bodies, 54, 332
*Isonandra gutta*, 474
Ispaghúl, 406
-ite, meaning of, 76
Ivory-black, 477

Jaborandi, 397
Jalap, Mexican male, 419
  resin, 419
  Tampico, 419
  true, 419
*Jalapa*, 419
*Jalapæ resina*, 419
  *resina*, impurities in, 618
Jalapic acid, 419
Jalapin, 419
Jalapinol, 419
Jaune brillant, 249
Jelly, 453
  vegetable, 405
Jervia, 396
Jervine, 396
Juices, 507
Juniper-oil, 465
Juniperis, 465

Kainit, 61
Kaladana resin, 420
Kali, 32
*Kalium*, 31
*Kamala*, 470
Kariyat, 400
Kelp, 273
Kermes mineral, 183
Kieserit, 116
Kilbride mineral, 151
Kiln, 106
Kilogram, 522, 523
Kilolitre, 528
Kilometre, 522
Kinate of quinia, 384
Kinetic theory, 24
King's blue, 477
Kino, 357
Kinone, 415
Kiwach, 241
Kokum butter, 458
Kola-nut, 398
Kosin, 470
Koussin, 470
Kousso, 470
*Kramericæ radix*, 357

Laboratory furniture, xii.
*Lac*, 451
Lac-dye, 476
Lactates, 346
Lactic acid, 346
Lactometer, 452
Lactose, 411
*Lactuca*, 400
Lactucin, 400
Lævogyrate, 410, 412
Lævoracemic acid, 320
Lævorotation, 412
Lævotartaric acid, 320
Lævulose, 410
Lakes, 139
Lampblack, 477
Lamps, gas-, 17, 23
*Lana philosophica*, 134
Lanthanum, 635
Lanthopine, 382
Larch bark, 357
Lard, 458
  benzoated, 458
  prepared, 458
*Laricis cortex*, 357
*Larix europæa*, 466
Larixin, 357
Larixinic acid, 357
Latent heat, 84
Laudanine, 382
Laudanosine, 382
Laughing-gas, 93
Laurate of glyceryl, 458
Laurel-camphor, 467
Lauric aldehyd, 466
Lauric acid, 458, 460
*Laurocerasi folia*, 414
Lavender-oil, 465
  -water, 465
Lavœsium, 635
Law, Avogadro's and Ampère's, 52
  Berthollet's, 378
  Boyle's, 52
  Charles's, 52
  of chemical combination by weight, 47 *et seq.*, 58
  by volume, 51 *et seq.*, 58
  concerning molecular weight, 54, 538

Law—
of constant proportions, 48, 510, 568
Dalton's, 47, 48
of diffusion, definition of, 57
Dulong and Petit's, 539
Gay Lussac's, 50
Graham's, 24
Henry and Dalton's, 85
Malaguti's, 378
Mariotte's, 52
multiple proportions, 48, 200, 290
reciprocal proportions, 48
solubility of gases in liquids, 85
Laws of chemical combination, 47, 52, 53
Lead, 209
acetate of, 210
analytical reactions of, 213, 485
antidotes to, 214
carbonate, 210
chloride, 213
chromate, 214
derivation of word, 32
gummate, 114
hydrato-carbonate, 210
in organic mixtures, detection of, 484
iodide, 212
nitrate, 212
oleate, 213
oxide, 210
oxyacetate, 211
oxychromate, 214
plaster, 213
puce-colored oxide or peroxide of, 212
pyrophorus, 158
quantitative estimation of, 546, 583
red, 212, 476
shot, 209
subacetate, 211
sugar of, 211
sulphate, 214
sulphide, 214
native, 209
test for, in water, 214
-tree, 215
volumetric estimation of solutions of acetate of, 549
Lead—
white, 210
Leadstone, 140
Leaf-green, 477
*Lecanora*, 477
Lees, 318
Legumin, 453
Lemon-chrome, 214
-juice, 324
-juice, estimation of mineral acid in, 590
-oil, 463
Length unit, 521
Lentisk tree, 470
Lepidolite, 227
Leptandrin, 400
Levisticum, 472
Levulose, 412
Lichen blue, 477
Lichenin, 404
Light carbonate of magnesium, 118
carburetted hydrogen, 433
magnesia, 119
Lignin, 407
Lime, caustic, 106
bisulphite, 306
chloride of, 113
-juice, estimation of mineral acid in, 590
-kiln, 106
-oil, 463
quick, 106
slaked, 107
superphosphate of, 327
-water, 107
Limestone, 105
magnesian, 117
mountain, 117
*Limonis cortex*, 463
*succus*, impurities in, 618
Limonite, 151
Line, 526
*Lini farina*, 459
*semina*, 459
Liniment of mercury, 195
Liniments, 507
*Linimentum ammoniæ*, 456
*calcis*, 457
Linoxin, 459
Linseed, 459
-cake, 459
oil, 459
tea, 406

Linum, 459
*usitatissimum*, 405
Liqueurs, 423
*Linuidambar orientale*, 473
Liquid camphor, 467
definition of, 57
Liquids, specific gravity of, 530
official, specific gravity of, 531
**Liquor** *ammoniæ*, 91
*ammoniæ*, impurities in, 618
*ammoniæ citratis*, 94
*ammoniæ fortior*, 91
*ammoniæ fortior*, impurities in, 618
*ammonii acetatis*, 92
*antimonii chloridi*, 180
*antimonii chloridi*, impurities in, 619
*antimonii chloridi*, estimation of antimony in, 580
*arsenicalis*, 167
*arsenicalis*, impurities in, 619
*arsenici chloridi*, 168
*arsenici et hydrargyri iodidi*, 166
*arsenici hydrochloricus*, 168
*arsenici hydrochloricus*, impurities in, 619
*atropiæ*, 393
*atropiæ sulphatis*, 393
*bismuthi et ammoniæ citratis*, estimation of bismuth in, 253
*bismuthi et ammoniæ citratis*, impurities in, 619
*calcis*, 107
*calcis*, impurities in, 619
*calcis chloratæ*, 114
*calcis chloratæ*, impurities in, 619
*calcis saccharatus*, 107
*calcis saccharatus*, impurities in, 619
*chlori*, 268
*chlori*, impurities in, 619
*ferri citratis*, 152
*ferri perchloridi*, 148
*ferri perchloridi fortior*, 148
*ferri perchloridi fortior*, impurities in, 619
*ferri perchloridi fortior*, estimation of iron in, 578
*ferri pernitratis*, 157
*ferri pernitratis*, impurities in, 619
**Liquor**—
*ferri pernitratis*, estimation of iron in, 578
*ferri persulphatis*, 153
*ferri persulphatis*, impurities in, 619
*ferri persulphatis*, estimation of iron in, 578
*ferri subsulphatis*, 149
*ferri tersulphatis*, 153
*gutta-perchæ*, 474
*hydrargyri nitratis acidus*, 199
*hydrargyri nitratis acidus*, impurities in, 619
*hydrargyri perchloridi*, 201
*iodi*, 274
*iodinii compositus*, 274
*lithiæ effervescens*, 227
*lithiæ effervescens*, impurities in, 619
*magnesiæ carbonatis*, 118
*magnesiæ carbonatis*, impurities in, 619
*magnesiæ citratis*, 119
*magnesii citratis*, 119
*morphiæ acetatis*, 382
*morphiæ hydrochloratis*, 382
*morphiæ sulphatis*, 382
*plumbi subacetatis*, 211
*plumbi subacetatis*, impurities in, 619
*plumbi subacetatis dilutus*, 211
*potassæ*, 62, 67
*potassæ*, impurities in, 619
*potassæ effervescens*, 72
*potassæ effervescens*, impurities in, 619
*potassæ*, neutralizing power of, 546
*potassæ*, specific gravity of, 532
*potassæ*, to prepare pure, 67
*potassii arsenitis*, 167
*potassii citratis*, 72
*potassii permanganatis*, 231
*sodæ*, 82
*sodæ*, impurities in, 619
*sodæ chloratæ*, 86
*sodæ chloratæ*, impurities in, 619
*sodæ chlorinatæ*, 86
*sodæ effervescens*, 85
*sodæ effervescens*, impurities in, 619

**Liquor—**
*sodii arseniatis*, 170
*strychniæ*, 390
*zinci chloridi*, 132
Liquorice, 412, 418
-sugar, 412
List of apparatus, xi.
of chemicals, xiii.
of reagents, xiii.
Litharge, 210
Lithates, 359
*Lithii carbonas*, 227
*carbonas*, impurities in, 619
*citras*, 227
*citras*, impurities in, 619
Lithic acid, 359
**Lithium**, 227
analytical reactions of, 228
carbonate of, 227
citrate of, 227
derivation of word, 33
flame, 228
fluoride of, 227
silicate of, 227
sulphate of, 227
urate of, 228
Litmus, 95, 477
paper, 95
solution of, 95
tincture of, 95
Litre, relation of, to pints, 522
Liver of sulphur, 68
Lixiviation, 87
Loadstone or lodestone, 140
*Lobelia*, 397
Lobelia vinegar, 298
Lobelina, 397
Lobeline, 397
Logwood, 29, 476
solution of bleached, by chlorine, 29
Long pepper, 397
Looking-glasses, 241
*Lotio hydrargyri flava*, 204
*hydrargyri nigra*, 204
Louisa-blue, 477
Lozenges, 507
Lucifers, 24
Lump sugar, 410
Lunar caustic, 218
Lupulin, 472
oleo-resin of, 472
Lupuline, 397
Lupulinic acid, 472
*Lupulus*, 472
Luteolin, 475
Luting, 97
fire-clay, 201
linseed meal, 97
Lycopodium, 459
impurities in, 619

Mace, 458
fixed oil of, 458
volatile oil of, 465
*Macis*, 458
Madder, 476
Magenta, 478
Magnesia, 119
effervescing citrate of, 120
impurities in, 619
calcined, 119
fluid, 118
hydrous carbonate of, 119
*Magnesia levis*, 119
*levis*, impurities in, 619
Magnesian limestone, 116
*Magnesii carbonas*, 118
*carbonas*, impurities in, 620
*carbonas levis*, 118
*carbonas levis*, impurities in, 620
*carbonatis, liquor*, 118
*sulphas*, 117
*sulphas*, impurities in, 620
Magnesite, 116
**Magnesium**, 116
analytical reactions of, 120
and ammonium, arseniate of, 120
and ammonium, phosphate of, 120
carbonate of, 120
chloride of, 116
citrate of, 120
derivation of word, 32
detection of, in presence of barium and calcium, 124
euxanthate of, 475
for analytical purposes, 174
limestone, 116
oxide of, 119
phosphate of, in bones, 111
purrate of, 475
quantitative estimation of, 575
separation from barium and calcium, 121, 124
silicate of, 352

**Magnesium**—
sulphate of, 117, 575
sulphite of, 306
Magnetic iron ore, 140
oxide of iron, 156
Magpie test for mercury, 208
Maize starch (fig.), 403
Malachite, 190
Malaguti's law, 378
Malate of atropine, 393
nicotine, 397
Malates, 347
Male fern oil, 459
Malic acid, 347
Mallow tea, 406
Malt, 405
substitute, 412
Maltose, 412
Manganate of potassium, 76, 231
**Manganese**, 230
analytical reactions of, 230
black oxide of, 230
Crum's test for, 233
derivation of word, 33
quantitative analysis of black oxide of, 576
sulphate of, 233
*Manganesii oxidum nigrum*, 230
*sulphas*, 233
Manganous chloride, 230
hydrate, 233
sulphide, 232
Mangosteen oil, 458
*Manihot* starch (fig.), 403
*Manna*, 413
impurities in, 620
Mannite, 413
Manufacturing chemists, 14
Manures, analysis of, 613
*Maronta* starch (fig.), 403
Maraschino, 423
Marble, 105
Margarine, 458
Margosa bark, 400
Marine soap, 458
Mariotte's law, 52
Marking-ink, 218
Marl, 137
*Marmor album*, 105
Marsh-gas, 433
Marshmallow, 405
Marsh's test for arsenicum, 173
Maryland senna, 415
Massicot, 210
Mastic, 470
*Mastiche*, 470
Mastichic acid, 470
Masticin, 470
Maté, 398
*Maticæ folia*, 400
*Matricaria chamomilla*, 463
Mauve, 478
May-apple, 395
Meadow-sweet, oil of, 442
Measurement of temperature, 513
Measures, 519 *et seq.*
Mechanical and chemical combination, difference between, 30, 35
Mechanical medicines, 241
Meconate of morphine, 381
Meconic acid, 348, 489
Meconidine, 382
Meconine, 382
Meconoisine, 382
Meerschaum, 352
*Mel*, 412
impurities in, 620
*boracis*, 333
*depuratum*, 412
*sodæ boratis*, 333
Melam, 355
Melasses, 412
Melegneta pepper, 465
Melissic acid, 460
Melting-points, Table of, 517
of fats, etc., to determine, 516
of metals, 517
Melissyl, palmitate of, 458
Melon-essence, 442
Memoranda, analytical, 259, 367
*Menispermum canadense*, 394
Mentha, 465
*Mentha piperita*, 465
*viridis*, 465
Menthene, 465
Menthol, 465
Mercuric ammonium, chloride of, 195, 206
chloride, 196, 200
cyanide, 280
hexiodide, 274
iodide, 196, 198
oxide, 203
oxynitrates, 199
oxysulphate, 200
nitrate, 198

**Mercuric—**
salts, 196
salts, analytical reactions of, 204, 484
sulphate, 199
sulphide, 206
*Mercurius vitæ*, 181
**Mercurious ammonium, chloride of, 206**
chloride, 196, 201
chromate, 208
iodide, 196
nitrate, 198
oxide, 204
salts, 195
salts, analytical reactions of, 207
sulphate, 199
sulphide, 207
**Mercury, 194**
amido-chloride of, 206
ammoniated, 205
ammonio-chloride of, 205
analytical reactions of, 204
antidotes to, 208
basic sulphate of, 200
black oxide of, 203
carbonates of, 208
cyanide of, 280
chlorides of, 200
derivation of word, 32
hexiodide, 274
iodides of, 196, 205
galvanic test for, 208
magpie test for, 208
native sulphide of, 194
nitrates of, 198
nomenclature of salts of, 195
of life, 181
in organic mixtures, detection of, 484
oxides of, 203
oxynitrates of, 199
oxysulphate of, 200
oxysulphide of, 207
quantitative estimation of, 581
subchloride of, 202
sulphates of, 199
sulphide of, 194, 206
yellow oxide of, 203
Meta, meaning of, 243
Metaboracic acid, 333
Metacinnamein, 473
Metallic elements, 15
Metalloids, 15
**Metals, 15**
of minor pharmaceutical importance, 226
quantitative estimation of, 567
Table of the fusibility of, 517
Metamerides, 407
Metamerism, 407
Metantimonic acid, 181
Metaphosphates, 348
Metaphosphoric acid, 348
Metastannates, 243
Metastannic acid, 243
Metastyrol, 474
Metathesis, 77, 292
Metavanadates, 332
Methenyl, 437
Methyl, 433
conia, 396
hydride of, 433
-protocatechuic aldehyde, 361
salicylate, 442
Methylamine, 379
Methylated spirit, 434
sweet spirit of nitre, 435
Methylene, dichloride of, 437
Methylic alcohol, 434
detected in presence of ethylic alcohol, 434
Metre, 522
relation of, to inches, 522, 527
Metric system, 520 *et seq.*
of weights and measures, its relation to the English, 522 *et seq.*
system, weights and measures of, 520 *et seq.*
*Meum*, 472
*Mezerei cortex*, 470
Mezereon, 470
*Mica*, 137
*Panis*, 411
Microcosmic salt, 371
Microscopic examinations of urinary sediments, 498
Microscopy of starches, 403
Microspectroscope, 482
Milk, 451
sugar, 411
sulphur, 304
Mimotannic acid, 357
Mineral acids, detection of, in organic mixtures, 486
chameleon, 231

Mineral—
kermes, 183
kilbride, 151
purple, 476
rouge, 476
Minerals, general analysis of, 369 *et seq.*
special analysis of, 369
Minim, 526
Minium, 210
Mint, 465
Mirbane, essence of, 443
*Mistura ferri aromatica*, 161
*ferri composita*, 144
*potassii citratis*, 325
Mixture, definition of, 57
different from chemical combination, 30
Mixtures, 507
Mohr's burette, 543
Moist sugar, 410
Molasses, 412
Molecular volume, 55
weight, 54, 266, 538
weights, definition of, 58
Molecule, definition of, 57
Molecules, 41, 57
Molybdenum, 635
sulphide, 331
Molybdic acid, 492
Monads, 122
Monamines, 380
*Monarda*, 466
Monobasic acids, 265
Monobasylous radicals, 265
Monobrom-camphor, 467
Mononitrocellulin, 408
Monsel's solution, 149
Mordants, 139
*Mori succus*, 476
Morphia or Morphine, 381
acetate of, 381
analytical reactions of, 382, 489
hydrochlorate of, 381
in organic mixtures, detection of, 489
quantitative estimation of, 607
*Morphiæ acetas*, 382
*hydrochloras*, 382
*hydrochloras*, impurities in, 620
*murias*, 382
*sulphas*, 382
Mosaic gold, 244
*Moschus*, 453
*moschiferus*, 453
Mother-liquor, 108
Motion from heat, 84
Mountain-blue, 477
limestone, 116
Mucic acid, 413
Mucilage of bael, 406
of gum acacia, 114
linseed, 405
marshmallow, 405
slippery elm, 406
starch, 404
squill, 406, 422
tragacanth, 114
*Mucilago acaciæ*, 114
*amyli*, 404
*tragacanthæ*, 114
*Mucuna pruriens*, 241
Mulberry calculus, 506
-essence, 442
-juice, 476
-sugar, 410
Mulder's process for estimating alcohol, 610
Multiple proportions, law of, 48, 197, 290
Murexid, 359
Musk, 453
deer, 453
Mustard, 447
artificial oil of, 447
essential oil of, 447
fixed oil of, 459
"plaster," 447
*Mylabris cichorii*, 468
*Myrcia acris*, 423
Myristate of glyceryl, 458
Myristic acid, 460, 465
*Myristica*, 465
Myristicene, 465
Myristicol, 465
Myristin, 458
Myronate of potassium, 447
Myrosin, 447
*Myroxylon Pereiræ*, 473
*Toluifera*, 473
Myrrh, 472
*Myrrha*, 472
Myrrhic acid, 473
*Myrtus communis*, 465

Naphthalic acid, 336
Naphthalin, 336

Narceia, 382
Narcotine. 382
*Narthex assafœtida*, 472
Natal aloes, 446
Nataloin, 446
*Natrium*, 31
Natural philosophy, 42
*Nectandra Rodiæi*, 394
*Nectandriæ cortex*, 394
Nectandria, 397
Needle iron ore, 151
Negative radicals, 443
Neptunium, 635
Neroli oil, 463
Nessler test, 548
Neutral chromate, 103
Neutralization, 93
**Nickel**, 235
  analytical reactions of, 235
  arsenio-sulphide of, 235
  cobalticyanide, 236
  cyanide, 236
  derivation of word, 34
  hydrate of, 236
  separation of, from cobalt, 236
  sulphide of, 236
Nickar nuts, 400
Nicotia, nicotine, nicotina, or nicotylia, 397
*Nicotina tabacum*, 397
Nieshnic bitters, 394
*Nihilum album*, 134
Nim, 400
Niobium, 636
**Nitrate** of ammonium, 93
  argent - ammon - ammonium, 206
  barium, 102
  bismuth, 250
  cadmium, 249
  iron, 157
  lead, 212
  mercury, 198
  potassium, 61, 287
  silver, 217
  silver, standard solution of, 556
  sodium, 81, 285
  strontium, 228
Nitrates, 285
  analytical reactions of, 288
  quantitative estimation of, 528
Nitre, 285
  cubic, 285
  sweet spirit of, 349, 430
Nitric acid, 287
  acid, antidotes to, 291
  acid in organic mixtures, detection of, 486
  acid, volumetric estimation of, 552
  anhydride, 287, 289
  anhydride, radical, and acid, in the liquid acid of different densities, Table showing the percentage of, 626
  oxide, preparation of, 289
  peroxide, 289
Nitrite of amyl, 442
  of ethyl, 349
  potassium, 349
**Nitrites**, 349
  analytical reactions of, 349
Nitrobenzol, 445
  in oil of bitter almonds, test for, 415
Nitrocellulin, 408
**Nitrogen**, 25
  derivation of word, 31
  in the atmosphere, 25
  oxides of, 289
  peroxide of, 289
  preparation of, 25
  properties of, 26
  quantitative estimation of, in organic compounds, 600 *et seq.*
  relative weight of, 26
Nitrohydrochloric acid, 186, 288
Nitromuriatic acid, 288
Nitrous acid, 289
  anhydride, 290
  ether, 349
  oxide, 290
Nomenclature of salts:—
  alkaloids, 379
  anhydrides, 84
  anhydrous bodies, 84
  -ate, 73
  bi, 72
  carbonization, evaporation, ignition, incineration, 101
  double salts, 79
  hydrates, 65
  hydrous bodies, 84
  hyper, 146

Nomenclature of salts—
-ic, -ous, 72, 76, 142
-ide, -ite, 76, 142
iron salts, 142
mercury compounds, 196
per, 149
Non-drying oils, 459
Non-metallic elements, 15
Non-metals, 15
Nordhausen sulphuric acid, 310
Notation, 41 *et seq.*
Notes, analytical, 259, 367
Nutmeg, expressed oil, 465
oil of, 465
Nutrition, plastic elements of, 453
Nux vomica, 391

Oak-bark, 356
black, 475
Oatmeal, 401
Occlusion, 247, 248
Ochre, 475
Octohedral, 172
Œnanthylic acid, 460
Official liquids, specific gravity of, 531
substances, volumetric estimation of, 543, 552, 555, 558, 561, 563
formula, 28
Officinal formula, 28
Oil, ajowan, 463
ajwain, 463
almond, 459
amber, 354
aniseed, 463
arachis, 459
apple, 441
benne, 460
bergamot, 463
bitter almond, 414
bitter almond, artificial, 443
boldo, 464
buchu, 464
cacao, 458
cajuput, 464
cake, 459
camphor, 467
cannabis indica, 469
capsicum, 395
caraway, 464
cardamoms, 464
cascarilla, 464
cassia, 464

Oil—
castor, 459
cedra, 463
chamomile, 463
cinnamon, 464
citron, 463
citronella, 464
cloves, 464
cocoa-nut, 458
cod-liver, 459
copaiva, 464
coriander, 464
croton, 459
cubeb, 464
cummin, 464
dill, 463
earth-nut, 459
eggs, 450
elder-flower, 466
eucalyptus, 465
fennel, 465
garcinia, 458
garlic, 448
geranium, 465
gingelly, 460
ginger, 467
ginger-grass, 465
grains of paradise, 465
ground-nut, 459
hop, 472
horsemint, 466
horseradish, 447, 463
Indian hemp, 469
jaborandi, 397
juniper, 465
lavender, 465
lemon, 463
lemon-grass, 467
lime, 463
linseed, 459
lycopodium, 459
mace, fixed, 458
mace, volatile, 465
male-fern, 459
mangosteen, 458
meadow-sweet, 442
melissa, 467
mustard, artificial, 448
mustard, fixed, 459
mustard, volatile, 466
myrtle, 465
neroli, 463
nutmeg, 465
nutmeg, fixed, 458

Oil—
nutmeg, volatile, 465
olibanum, 473
olive, 459
omum, 463
orange-flower, 463
orange-rind, 463
origanum, 465
orris, 465
pennyroyal, 465
pepper, 397
peppermint, 465
petit grain, 463
pilocarpine, 397
pimento, 464
ptychotis, 463
rose, 465
rosemary, 465
rue, 465
saffron, 475
sandal-wood, 466
sassafras, 466
savin, 466
sesame, 460
shark liver, 460
spearmint, 465
sperm, 458
star-anise, 463
sweet flag, 466
teel, 460
theobroma, 458
thyme, 466
turmeric, 466
turpentine, 466
valerian, 467
verbena, 467
vitriol, 389
water-hemlock, 464
wine, 447
winter-green, 442
wormseed, 467
Oils and fats, composition of, 456
Oils, analysis of, 613
drying, 459
essential, 461
tested for alcohol, 462
fixed, 459
non-drying, 459
volatile, 461
volatile, process for, 462
Ointments, 507
Okra, 406
Olea destillata, impurities in, 620
Oleate of glyceryl, 456
of lead, 213
Oleates, 456
Olefiant gas, 447
Oleic acid, 456
Oleine, 456
Oleo-resins, 470
*Oleoresina capsica*, 472
*cubebæ*, 472
*lupulinæ*, 472
*piperis*, 472
*zingiberis*, 472
*Oleum amygdalæ amaræ*, 414
*amygdalæ dulcis*, 459
*andropogi citratis*, 467
*anethi*, 463
*anisi*, 463
*anthemidis*, 463
*arachis*, 459
*bergamii*, 463
*cajuputi*, 464
*camphoræ*, 467
*carui*, 464
*caryophylli*, 464
*cinnamomi*, 464
*copaibæ*, 464
*coriandri*, 464
*crotonis*, 459
*cubebæ*, 464
*ethereum*, 447
*fœniculi*, 465
*gaultheriæ*, 442
*juniperi*, 465
*lavandulæ*, 465
*limonis*, 463
*lini*, 459
*macis*, 458, 465
*menthæ piperitæ*, 465
*menthæ viridis*, 465
*morrhuæ*, 459
*myristicæ*, 465
*myristicæ expressum*, 458
*myrti*, 465
*olivæ*, 459
*origani*, 465
*phosphoratum*, 328
*pimentæ*, 464
*pulegii*, 465
*ricini*, 459
*rosæ*, 465
*rosmarini*, 465
*rutæ*, 465
*sabinæ*, 466
*sassafras*, 466

*Oleum*—
*sinapis*, 466
*succini*, 354
*succini rectificatum*, 354
*terebinthinæ*, 466
*theobromæ*, 458
*thymi*, 466
*tiglii*, 459
*valerianæ*, 467
Olibanum, 463
Olive-oil, 459
Omentum, 458
Omum oil, 463
Opal, 352
*Ophelia chirata*, 350
Ophelic acid, 350
Opianic acid, 382
Opianine, 382
Opium, 382
detection of, in organic mixtures, 489
estimation of morphia in, 607
impurities in, 620
vinegar, 297
Orange-chrome, 214
Orange-flower, 463
-flower-oil, 463
-rind oil, 463
-wine, 423
Orchil, 477
Orchis tuber, 406
Orcin, 477
Ordeal-poison, 397
Orellin, 476
Organic analysis, 596
bases, 379
chemistry, 378
compounds, 378
Orpiment, 475
Orris, butter of, 465
oil of, 465
Ortho, meaning of, 348
Orthophosphates, 348
Orthophosphoric acid, 348
Orthovanadates, 332
*Oryza sativa*, 401
*Oryza*, starch of (fig.), 403
*Os*, 111
*Os ustum*, 111
Osmium, 248, 636
Otto of rose, 465
Ounce, 519
Ourari, 391
-ous, meaning of, 76, 141
*Ovi vitellus*, 450
*Ovum*, 450
Ox-bile, 457
-gall, 457
Oxalate of ammonium, 95
barium, 317
calcium, 317
cerium, 229
iron, 316
potassium, 316
silver, 317
sodium, 316
strontium, 229
**Oxalates**, 316
analytical reactions of, 317, 486
from phosphates and ferric oxide, separation of, 373
quantitative estimation of, 592
Oxalic acid, 316
acid, antidotes, 317
acid, chemically pure, 316
acid, in organic mixtures, detection of, 486
acid, standard solution of, 546
**Oxide** of aluminium, 139
antimony, 181
bismuth, 252
calcium, 105
chromium, 238
copper, 191
iron, black, 156
lead, 210
magnesium, 119
manganese, 230
mercury, 203
silicon, 352
silver, 218
tin, 242
zinc, 133
Hubbuck's, 134
Oxides identified, 375
Oxides of nitrogen, 290
Oxidizing flame, 371
Oxyacetate of copper, 192
of lead, 211
Oxyacid salts, 287
Oxyacids of sulphur, 345
Oxycarbonate of bismuth, 252
Oxychloride of antimony, 180
Oxychromate of lead, 214
**Oxygen**, 16
derivation of word, 31

**Oxygen—**
from ozone and antozone, 271
in the air, 16, 26
its relation to animal and vegetable life, 19
preparation of, 16
properties of, 20
quantitative estimation of, in organic compounds, 597 *et seq.*
solubility in water, 19
specific gravity of, 24
weight of 100 cubic inches, 538
Oxygenated water, 103
Oxyhydrates of iron, 152
Oxyiodate of iron, 296
Oxymel, 412
of squill, 412
Oxynitrates of mercury, 203
bismuth, 252, 253
Oxysalts, 287
Oxysulphate of iron, 143
mercury, 200
Oxysulphide of antimony, 182
mercury, 207
Ozokerite, 458
Ozone, 275, 502

Palas tree, 357
Palladium, 248, 636
Palm-oil, 458
Palmitate of cetyl, 458
glyceryl, 458
melissyl, 458
Palmitic acid, 460
Palmitine, 458
Pancreatin, 455
*Papaver rhœas*, 476
*somniferum*, 381
*Papaveris capsulæ*, 381
Papaverine, 382
Paper, bibulous, 108
for filtering, 108, 568
Papers, test-, 95
Para, meaning of, 320
Paracyanogen, 280
Paraguay tea, 398
Parallin, 421
Paratartaric acid, 320
*Pareiræ radix*, 394
Paricine, 394
Parietinic acid, 337
Parigenin, 421
Parilla, 394
Paris blue, 477
red, 477
Particles, elementary, 46
Patent sugar, 412
Pear wine, 423
Pearlash, 61
Pearl-barley, 401
Pearl-sago starch (fig.), 403
-white, 252, 478
Peas, 453
Pectin, 405
Pelargonic acid, 460
Pelletierine, 357
Pellitory root, 470
Pelosia, 394
Pelosine, 394
Pentachloride of antimony, 180
Pentathionic acid, 345
Pepper, black, 397
cayenne, 395
cubeb, 397
long, 397
white, 397
oil of, 397
resin of, 470
Peppermint oil, 465
Pepsin, 454
Peptone, 455
Per-, meaning of, 146
Perbromates, 271
Percha tree, 474
Perchlorate of potassium, 294
Perchloric acid, 294
Perchloride of gold, 245
iron, 146
platinum, 247
tin, 242
Perfumes, 463
Perhydrate of iron, 150
Periodide of ammonium, 274
mercury, 274
potassium, 274
Permanganate of potassium, 76, 231
its use in volumetric analysis, 558
Pernitrate of iron, 157
Peroxide of barium, 103
hydrogen, 103
iron, 151
iron, hydrated, 150
lead, 212
nitrogen, 289

Peroxyhydrate of iron, 150
Persian berries, 475
Persulphate of iron, 149
Persulphide of hydrogen, 303
Peru, balsam of, 473
Peruvine, 473
Petalite, 227
Pettenkofer's test for presence of bile, 457
*Peumus boldus*, 464
Pewter, 179, 210, 241
Phæoretine, 337
Pharaoh's serpents, 355
*Pharbitisin*, 420
*Pharbitis nil*, 420
Pharmaceutical chemists, 14
Pharmaceutical Society of Great Britain, examinations of, 14
Pharmacists, 14
Pharmacy, 14
Phenic acid, 444
  alcohol, 444
Phenol, 444
Phenyl, 444
**Phosphate** of ammonium, 94
  barium, 331
  calcium, 105, 110, 331
  iron, 144, 331
  magnesium and ammonium, 120
  magnesium and ammonium from oxalates and ferric oxides, separation of, 373
  magnesium, in bones, 111
  silver, 219
  sodium, 112
  sodium, how prepared from phosphate of calcium, 112
Phosphates, 327
  analytical reactions of, 330
  quantitative estimation of, 593
**Phosphites**, 350
  test for, 351
Phosphomolybdic acid, 492
Phosphoretted hydrogen, 343
  oil, 328
Phosphoric acid, 24, 329, 350
  acid, diluted, 329
  acid, quantitative estimation of free, 593
  acid, table of strengths of solutions of, 630
  anhydride, 25, 348
Phosphorous acid, 350
**Phosphorus**, 24, 327
  combustion of, 24
  derivation of word, 31
  detection of, in organic mixtures, 487
  granulated, 328
  pill, 328
  properties of, 24
  red or amorphous, 328
  trihydride, 343
Phthalic acid, 336
Phyllocyanin, 477
Phylloxanthin, 477
Physical isomerides, 407
Physics, 42
*Physostigmatis faba*, 397
Physostigma, 397
Physostigmine, 397
Picric acid, 445
Picrotoxin, 420
Pigments, 475
*Pigmentum nigrum*, 477
Pills, 507
Pilocarpine, 397
*Pilula ferri carbonatis*, 144
  *ferri iodidi*, 31
  *hydrargyri*, 195
  *hydrargyri subchloridi composita*, 202
  *phosphori*, 328
  *plumbi cum opio*, 211
  *quiniæ*, 385
*Pilulæ antimonii compositæ*, 202
  *ferri compositæ*, 144
Pimaric acid, 469
Pimento, 464
  oil, 464
*Pimpinella anisum*, 463
Pine-apple, essence of, 441
Pinic acid, 469
Pink saucers, 476
  the common, 421
Pins, 241
Pint, 526
*Pinus*, 466, 469
*Piper nigrum*, 397
Piperia, 397
Piperic acid, 398
Piperidia, 398
Piperidine, 398
Piperine, 397
*Pistachia terebinthus*, 466
Pitch, 471

Pitch—
Burgundy, 470
*Pix burgundica*, 470
*canadensis*, 471
*liquida*, 471
*Plantago ispaghula*, 406
Plants and animals, complementary action on air, 19
Plaster of ammoniacum and mercury, 195
mercury, 195
Paris, 105, 478
Plasters, 213, 507
Plastic elements of nutrition, 453
Platinic salts, 247
Platinous salts, 247
**Platinum**, 246
analytical reactions of, 247
and ammonium, chloride of, 98, 572
and potassium, chloride of, 78
black, 247
derivation of word, 34
foil, 76, 246
perchloride of, 247
residues, to recover, 248
spongy, 248
sulphide of, 247
Pleurisy-root, 400
Plumbago, 30
*Plumbi acetas*, 211
*acetas*, impurities in, 620
*carbonas*, 210
*carbonas*, impurities in, 620
*emplastrum*, 213
*iodidum*, 212
*nitras*, 212
*oxidum*, 210
*oxidum*, impurities in, 620
*subacetatis, liquor*, 211
"Plummer's pills," 202
Plumbic peroxide, 212
acetate, sulphate, etc. (*vide* salts of lead).
*Plumbum*, 32
*Pocula emetica*, 179
*Podophylli radix*, 395
*resina*, 395
Poisonous alkaloids, 488, 490
Poisons, antidotes to (*vide* Antidotes, detection of, in organic mixtures, 483 *et seq.*).
Polybasic acids, 265
Polybasylous radicals, 265
Polychroite, 475
*Polygala senega*, 421
Polygalic acid, 421
Polymerism, 406
Polymorphism, 407
Polymorphous bodies, 407
Polysulphide of calcium, 303
Pomegranate rind, 357
Pomegranate-root bark, 357
Porcelain, 353
Port wine, 423
Porter, 423
Portland cement, 353
Positive radicals, 443
Potash alum, 138
solution of caustic, 67
solution of caustic, to prepare pure, 67
sulphurated, 68
table of strength of solutions of, 633
vol. estim. of sol. of, 546
-water, 67
Potashes, 61
Potassa, 67
*Potassa caustica*, 67
*caustica*, impurities in, 620
*sulphurata*, 68
*sulphurata*, impurities in, 620
*Potassæ* (vide *Potassii*).
*effervescens, liquor*, 72
*liquor*, 62, 67
*liquor*, to prepare pure, 67
*prussias flavæ*, 280, 339
Potassic hydrate, etc. (*vide* salts of potassium).
*Potassii acetas*, 69
*acetas*, impurities in, 620
*bicarbonas*, 71
*bicarbonas*, impurities in, 620
*bichromas*, 237
*bitartras*, 61, 73, 79, 319
*bromidum*, 76
*bromidum*, impurities in, 620
*carbonas*, 61
*carbonas pura*, 61
*carbonas impura*, 61
*carbonas*, impurities in, 620
*chloras*, 293
*chloras*, impurities in, 620
*citras*, 72
*citras*, impurities in, 620
*cyanidum*, 279
*et sodii tartras*, 73, 85

*Potassii*—
*ferridcyanidum*, impurities in, 620
*ferrocyanidum*, 280
*hypophosphis*, 343
*iodidum*, 74
*iodidum*, impurities in, 620
*nitras*, 73, 285
*nitras*, impurities in, 620
*permanganas*, 76, 231
*permanganas*, impurities in, 620
*sulphas*, 73, 287
*sulphas*, impurities in, 620
*sulphuretum*, 68
*tartras*, 73
*tartras*, impurities in, 620
*tartras acida*, 78, 319
*tartras acida*, impurities in, 620
Potassio-citrate of iron, 152
-tartrate of antimony, 182, 319
-tartrate of iron, 152
**Potassium**, 61
acetate of, 69
acid carbonate of, *vide* bicarbonate.
acid tartrate of, 79
analytical reactions of, 77
antimoniate of, 88
bicarbonate of, 71, 547
chemically pure, 552
bichromate of, 237
bitartrate of, 79, 319
borotartrate of, 334
bromate of, 76
bromide of, 76
carbonate of, 61, 547
chemically pure, 552
chlorate of, 16, 293
chloride of, 78
chromate of, 103
and platinum, chloride of, 78
citrate of, 72
cobalticyanide of, 235
cyanate of, 337
cyanide of, 279
derivation of word, 31
ferrate of, 141
ferridcyanide of, 340
ferrocyanide of, 280, 339
-flame-test, 79
hydrate of, 62
hydrate, to prepare pure solution, 67
**Potassium**—
hypophosphite of, **343**
iodate of, 75, 296
iodide of, 73
manganate of, 76, **231**
myronate of, 447
nitrate of, 61, 73, **285**
oleate of, 456
perchlorate of, **294**
periodide, 274
permanganate of, 76, **231**
preparation of, 61
properties of, 61
quantitative estimation of, 544, 546
quantivalence of, 61
red chromate of, 103, **237**
red prussiate of, 340
and sodium, tartrate of, **85**
salts, analogy of, to sodium salts, 87
sodium, and ammonium, separation of, 100
sources of, 61
sulphate of, 73, 287
sulphides of, 68
sulphocyanate of, **354**
sulphurated, 68
tartrate of, 73
tartrate of, acid, 61, 78, **319**
tri-iodide, 274
yellow chromate of, 103
yellow prussiate of, 280, **339**
Potato-oil, 441
starch (fig.), 403
Poultices, 507
Pound, 526
Powder, bleaching-, 113
Powders, 507
soda-, 87
specific gravity of, 534
Practical analysis, 99
Precipitant, 78
Precipitate, 78
Precipitated chalk, 108
sulphur, 303
Precipitates, soluble, in solutions of salts, 282
to wash, 108, 109, 569
to weigh, 568, 570, 573
Precipitation, 78
Preparations of the British Pharmacopœia, chemical, 509
galenical, 507

Prepared carbonate of calcium, 110
chalk, 110
lard, 458
suet, 458
Pressure, correction of vol. of gas for, 536
gauges, 512
Prickly ash, 394
Principles of Chemical Philosophy, 36 (*see* also "Laws").
Printer's ink, 477
Prismatic nitre, 285
Prophetin, 417
Proportions, atomic, 47, 197
constant, 47, 58
multiple, 48, 58
reciprocal, 48, 58
Proof spirit, 423
Propionic acid, 460
Propyl, 433
Propylamine, 380
Protocatechuic aldehyd, 361
Protopine, 382
Proximate analysis, 596
Prune, 412
*Prunum*, 412
*Prunus serotina*, 415
*virginiana*, 415
Prussian blue, 340, 477
Prussiate of potash, red, 341
of potash, yellow, 279, 339
Prussic acid, 279
Pseudaconitine, 392
Pseudomorphia, 382
*Pterocarpi lignum*, 476
*Pterocarpus santilinus*, 476
*Ptychotis ajowan*, 463
Ptyalin, 503
Puce-colored oxide of lead, 212
Puddling, iron, 141
*Pulvis algarothi*, 181
*angelicus*, 181
*antimonialis*, 183
*ipecacuanhæ compositus*, 396
*Pulveres effervescentes*, 87
*aperientes*, 322
*Pumice stone*, 352
*Punica granatum*, 359
Purified ox-bile, 457
Purple of Cassius, 246
foxglove, active principle in, 416
Purpurine, 497
Purrate of magnesium, 475
Purree, 475
Pus, in urine, 502
Putty-powder, 243
Pyrethrin, 470
*Pyrethri radix*, 470
*Pyrethrum carneum*, 470
Pyrites, copper, 190
iron, 140
Pyroarseniate of sodium, 169
Pyroarseniates, 170
Pyrogallic acid, 358
acid, use of, in gas analysis, 358
Pyroligneous acid, 297
Pyrolusite, 230
Pyrometers, 516
Pyromorphite, 332
Pyrophorus, 159
Pyrophosphates, 332, 351
Pyrophosphoric acid, 349, 351
Pyrovanadates, 332
Pyroxylic spirit, 434
*Pyroxylin*, 408
Pyroxylin, 408

Quadrivalence, 56
Qualitative analysis, 99
Quantitative analysis, 510 *et seq.*
Quantivalence, 55, 121
of atoms, definition of, 58
of acidulous radicals, 66
Quartz, 352
*Quassiæ lignum*, 400
Quassin, 400
Quercitrin, 475
Quercitron, 475
*Quercus cortex*, 356
*tinctoria*, 475
Quevenne's iron, 158
Quicklime, 106
Quinamia, 388
Quinamine, 388, 389
Quince seeds, 406
*Quiniæ sulphas*, 384
*sulphas*, impurities in, 621
*valerianas*, 385
Quinia, or quinine, 384
amorphous, 605
analytical reactions of, 386
citrate, 386
citrate of iron and, 153
De Vrij's process for estimating, 603
disulphate of, 385

**Quinia—**
iodosulphate, 387
kinate of, 384
quantitative estimation of, 603
sulphate of, 385
Quinia wine, 385
Quinicia, 388
Quinicine, 388
Quinidia, 388
Quinidine, 388, 389
Quinine, 384, 388, 389
Quiniretin, 388
Quinquivalence, 56

**Radicals,** 66
acidulous, 60, 263
acidulous, formulæ of, 122
alcohol, 433
basylous, 60, 122
definition of, 60
negative, 443
positive, 443
Racemic acid, 320
Rai, 447
Raisins, 412
Raspberry, sugar in, 410
Ratafia, 423
Rational formulæ, 424
Reactions, analytical, 62
synthetical, 62
Reagents, list of, xv.
Real alcohol, 425
Realgar, 167
Reaumur's thermometer, 514
Reciprocal proportions, law of, 50
Rectification, 126
Rectified oil of turpentine, 466
spirit, 126, 423
Red chromate of potassium, 237
coloring-matters, 475
corpuscles in blood, 501
earth, 476
enamel colors, 477
gravel, 497
hæmatite, 140
iodide of mercury, 196
litmus-paper, 95
lead, 212, 476
ochre, 476
oxide of iron, 150, 476
phosphorus, 328
-poppy petals, 476
Red—
precipitate, 203
prussiate of potash, 340
-rose petals, 476
sandal-wood, 476
sanders-wood, 476
Venetian, 152
Reduced indigo, 290
iron, 157
Reinsch's test for arsenicum, 192
Relations of gases, liquids, and solids, 42
Relative weight of hydrogen and oxygen, 24
Rennet, 451
*Reseda luteola*, 475
Resin, 469
of arnica, 469
of cannabis, 469
of capsicum, 469
of castor, 469
of copal, 469
of dragon's blood, 469
of ergot, 470
of guaiacum, 418, 470
of Indian hemp, 469
of jalap, 419, 470
of kamala, 470
of kousso, 470
of mastic, 470
of mezereon, 470
of pepper, 397
of podophyllum, 395
of pyrethrum, 470
of rottlera, 470
of scammony, 421
*Resina*, 469
*jalapæ*, 419
*podophylli*, 395
*scammoniæ*, 421
Resinoid substances, 469
Resins, 469
Respiratory materials of food, 453
Retort, 126
*Rhamni succus*, 416, 477
Rhamnin, 475
*Rhamnus catharticus*, 477
Rhaponticin, 337
Rhatany root, 357
*Rhei radix*, 337
*radix*, impurities in, 621
Rheic acid, 337
Rhein, 337
Rheumin, 337

Rhœadin, 382
*Rhœados petala*, 476
Rhodium, 248
Rhubarb, oxalate of calcium from, 500
Rhubarbic acid, 337
Rhubarbarin, 337
*Rhus cotinus*, 475
Rice, 401
Rice-starch (fig.), 403
Ricinine, 459
Ricinoleate of glyceryl, 459
Ricinoleine, 459
Ringworm powder, 337
*Roccella*, 477
Rochelle salt, 85, 320
Rock-salt, 81
Roll sulphur, 302
Roman cement, 353
*Rosæ caninæ fructus*, 412
*centifoliæ petala*, 465, 476
*gallicæ petala*, 476
Roscoe's vanadium, 332
Rosaniline, 478
Rose-oil, 465
petals, 465, 476
-water, 462
Rosemary-oil, 465
Rosin, 469
Rotten stone, 137
*Rottlera tinctoria*, 470
Rottlerin, 470
Rouge, animal, 337, 476
mineral, 151, 476
vegetable, 476
*Rubia tinctorum*, 476
Rubidium, 636
Ruby, 137
Rue-oil, 465
Rum, 423
Rumex, 337
Rumicin, 337
Rust of iron, 141
Rutate of glyceryl, 458
Ruthenium, 248
Rutic acid, 458
aldehyd, 466

*Sabadilla*, or sabadilline, 399
*Sabinæ, oleum*, 466
Saccharated carbonate of iron, 143
carbonate of iron, volumetric estimation of, 562
Saccharic acid, 413
Saccharimetry, 608
"Saccharine," 412
Saccharine substances, 401, 409
Saccharometer, 609
*Saccharum lactis*, 411
*purificatum*, 410
*ustum*, 412
Safety-lamp, 23
Safety-tube, 267
Safflower, 476
Saffranin, 475
Saffron, 475
bastard, 426
dyer's, 476
Safren, 466
Safrol, 466
Sago, 401
Sago starch (fig.), 403
Sal ammoniac, 90
Salep, 406
Salicin, 420
Salicyl, hydride of, 420, 442
Salicylate of methyl, 442
Salicylic acid, 442, 445
Salicylous acid, 442
Saligenin, 420
Saliva, 503
*Sal prunella*, 286
Salseparin, 421
Salt, common, 81
definition of a, 61
of sorrel, 316
Saltpetre, 285
Chili, 285
**Salts**, acid, 302
action of the blowpipe on, 370
action of heat on, 370
action of sulphuric acid on, 371
of ammonium, volatility of, 99
analogies of, 87
analysis of insoluble, 369
constitution of, 62, 125, 263, 286, 300, 377
formation of, 70
nomenclature of, 72, 75
of iron, nomenclature of, 141
physical properties of, 370
substitution of for each other, 87
table of the solubility or insolubility of, in water, 365
*Sal volatile*, 93
Sambucene, 466

*Sambuci flores*, 466
Sand, 352
-bath, 28
-tray, 28, 201
Sandal-wood, oil of, 466
red, 466, 476
white, 466
yellow, 466
Sandstone, 352
Sanguinaria, 398
vinegar, 298
Sanguinarina, 398
Santalin, 476
*Santalum album*, 466
*Santonica*, 420
Santonic acid, 420
Santonin, 420
*Santoninum*, 420
impurities in, 621
Santoniretin, 420
Sapan-wood, 476
Sap-green, 477
*Sapo animalis*, 457
*durus*, 456
*durus*, impurities in, 621
*mollis*, 456
*mollis*, impurities in, 621
*viridis*, 457
Saponin, 421
Sapphire, 137
*Sarcinæ ventriculi*, in urine, 504
Sarcolactic acid, 347
Sarsaparilla, 421
*Sarzæ radix*, 421
Sassafras-oil, 466
*Sassafras radix*, 466
Sassafrol, 466
Saturated solutions, boiling-points of, 516
Saturating power of citric acid, 325, 623
power of tartaric acid, 321, 623
Saturation, 69
tables, 321, 325, 623
Saturn, 210
Saturnine colic, 210
Savin oil, 466
Saxon blue, 477
Saxony blue, 477
Scale compounds of iron, 152
*Scammoniæ radix*, 421
*resina*, 421
*resina*, impurities in, 621
Scammonin, 421
Scammoniol, 421
*Scammonium*, 421
Scammony, resin of, 421
Scents, 463
Scheele's green, 177
Schist, 137
Schönbein's test for hydrocyanic acid, 284
Schweinfurth green, 177
Science of chemistry, 14
*Scilla*, 422
Scilletin, 422
Scillitin, 422
Sclerotic acid, 470
Sclerotinic acid, 470
*Scoparii cacumina*, 398
Scoparin, 398
Sea-salt, 81
Sediments, urinary, 496
urinary, microscopic examinations of, 498
Seidlitz powder, 321
Selenion, 636
Selenium, 303, 636
*Senegæ radix*, 421
*Senna alexandrina*, 415
*indica*, 415
Sepia, 477
Serolin, 451
*Serpentariæ radix*, 400
Serpent's excrement, 359
Sesame oil, 460
*Sesame indicum*, 460
*Sevum præparatum*, 458
Sexivalence, 56
Shale, 137
Shark-liver oil, 460
Shellac, 476
Sherry-wine, 423
Sienna, 477
Sifting, an aid to an analysis, 370
Silica, 352
Silicate of aluminium, 137
calcium, 105, 353
magnesium, 353
**Silicates**, 353
quantitative estimation of, 595
tests for, 353
Silicic acid, 353
anhydride, 353
Siliciuretted hydrogen, 353
Silico-fluoride of barium, 104

Silicon, chloride of, 354
derivation of word, 33
fluoride of, 354
hydride of, 353
oxide of, 353
**Silver**, 216
ammonio-nitrate of, 177, 206
analytical reactions of, 218
antidotes to nitrate of, 219
arseniate of, 177, 219
arsenite of, 177
bromide of, 219
chloride of, 217
chromate of, 219
citrate of, 326
coinage, 216
cyanide of, 219
by cupellation, estimation of, 584
derivation of word, 32
extraction of, 216
German, 130
iodide of, 219
nitrate of, 216, 217
oxalate of, 317
oxide of, 218
phosphate of, 219
pure, 217
quantitative estimation of, 584
standard solution of nitrate of, 556
sulphide of, 219
sulphite of, 307
tartrate of, 322
tree, 219
volumetric estimation of, 584
Sinalbin, 447
Sinapine, 447
*Sinapis*, 447
impurities in, 621
Sinigrin, 447
Siphon, *vide* Syphon.
Size, 454
Slaked lime, 107
Slate, 137
Smalt, 234, 477
Smilacin, 421
Soap, ammonium, calcium, green, hard, potassium, sodium, soft, 456, 457
Soap-curd, 457
Soap-stone, 478
-wort, 421
Socaloin, 446
Socatrine aloes, 446
"Soda," 313, 550
Soda, solution, table of strength of, 633
Soda-alum, 138
-ash, 87, 550
caustic, 81
*Soda*, 82
*caustica*, 82
*caustica*, impurities in, 621
*tartarata*, 85
*tartarata*, impurities in, 621
Soda-lime, 600
-powders, 87
solution of chlorinated, 86, 566
standard solution of, 553
valerianate of, 360
volumetric estimation of, 547
water, 85
Sodæ, *vide* Sodii.
*Sodii acetas*, 82
*acetas*, impurities in, 621
*arsenias*, 170
*arsenias*, impurities in, 621
*bicarbonas*, 83
*bicarbonas*, impurities in, 621
*boras*, 333
*carbonas*, 84, 313
*carbonas*, impurities in, 621
*carbonas*, *exsiccata*, 84
*chloratæ*, *liquor*, 86, 566
*chlorinatæ*, *liquor*, 86
*chloridum*, 81
*citro-tartras effervescens*, 86
*hypophosphis*, 343
*hyposulphis*, impurities in, 621
*liquor*, 82
*nitras*, 285
*nitras*, impurities in, 621
*phosphas*, 112
*phosphas*, impurities in, 621
*sulphas*, 268
*sulphas*, impurities in, 621
*sulphis*, 306
*valerianas*, 360
*valerianas*, impurities in, 621
Sodic carbonate, etc., *vide* salts of sodium.
Sodio-citrate of iron, 152
-tartrate of iron, 152
**Sodium**, 81
acetate of, 82

**Sodium—**
acid carbonate of, 83, 547
acid sulphate of, 268
acid tartrate of, 79
analytical reactions of, 88
and aluminium, double chloride of, 137
antimoniate of, 88
arseniate of, 170
arsenite of, 169
bicarbonate of, 83
chemically pure, 551
bisulphite of, 306
bromate of, 87, 271
bromide of, 87, 271
carbonate of, 81, 87, 313, 547
chemically pure, 551
carbonate of, manufacture of, 87, 313
chlorate of, 87
chloride of, 81
cholate, 457
citrate of, 87
derivation of word, 31
-flame, 88
glycocholate, 457
hydrate of, 81
hypochlorite of, 86
hypophosphite of, 343
hyposulphite of, 344
iodate of, 87
iodide of, 87
manganate of, 87
nitrate of, 81, 285
other compounds of, 87
oxalate of, 316
permanganate of, 87
phosphate of, 112
phosphate of, how prepared from phosphate of calcium, 112
potassium and ammonium, separation of, 100
pyroarseniate of, 169
quantitative estimation of, 547, 571
salts, analogy of, to potassium salts, 87
salts, sources of, 81
sulphate of, 268
sulphite of, 306
taurocholate, 458
valerianate of, 360
Soft soap, 456
Soils, analysis of, 613
Solania, 398
Solanidine, 398
Solanine, 398
*Solanum dulcamara*, 398
*tuberosum*, starch of (fig.), 403
Solazzi juice, 418
Solder, 209, 241
Solid, definition of, 57
fats, 456
potash, 67
Solids lighter than water, to take the specific gravity of, 535
to take the specific gravity of, 533 *et seq.*
Solubility of carbonic acid gas in water, 85
of gases in liquids, 85
of precipitates in strong solutions of salts, 282
or insolubility of salts in water, Table of, 365
Soluble cream of tartar, 334
glass, 353
substances, to take the specific gravity of, 535
tartar, 73
**Solution** of acetate of ammonium, 92
acetate of potassium, 69
acetate of sodium, 82
albumen, 449
ammonia, 91
ammonio-nitrate of silver, 177, 206
ammonio-sulphate of copper, 177, 206
ammonio-sulphate of magnesium, 120, 593
arsenic in acid, 168
arsenic in alkali, 167
boracic acid, 333
bromine, 275
carbonate of ammonium, 92
carbonate of magnesium, 119
chloride of ammonium, 90
chloride of antimony, 180
chloride of barium, 103
chloride of calcium, 105
chloride of calcium, saturated, 106
chloride of gold, 245
chloride of tin, 242
chloride of zinc, 132

**Solution of—**
chlorinated lime, 566
chlorinated soda, 86, 566
chlorine, 28
citrate of ammonium, 94
citrate of magnesium, 119
ferridcyanide of potassium, 340
ferrocyanide of potassium, 339
gelatine, 453
iodate of potassium, 75, 296
iodide of potassium, 75
iodine, 273
lime, 107
litmus, 95
nitrate of mercury, 199
oxalate of ammonium, 95
perchloride of iron, 148
perchloride of mercury, 201
perchloride of platinum, 247
pernitrate of iron, 157
persulphate of iron, 149
phosphate of sodium, 112
phosphoric acid, 329
potash, 62, 67
red prussiate of potash, 340
soda, 82
strychnia, 390
subacetate of lead, 211
sulphate of calcium, 114
sulphate of indigo, 290
sulphate of iron, 142
sulphide of ammonium, 96
sulphydrate of ammonium, 95
tartaric acid, 320
yellow prussiate of potash, 339
Sonnenschien's process for poisonous alkaloids, 491
Soot, 30
*Soubresants*, 281
Source of heat, 17
Sovereign, weight of the, 245
*Soymida febrifuga*, 400
Spanish liquorice, 418
Spar, fluor, 341
heavy, 102
Sparteia or sparteine, 398
Spathic iron ore, 141
Spearmint oil, 465
Specific gravity, 529
gravity of gases, 536
gravity of liquids, 530
Specific—
gravity of official liquids, 531
gravity of oxygen, 24
gravity of powders, 534
gravity of solids, 533
gravity of solids lighter than water, 535
gravity of soluble substances, 535
Specific heat, 128
weight, 529
Spectroscope, 482
Spectrum analysis, 482
Specular iron ore, 141
Speculum metal, 241
Speiss, 235
Spermaceti, 458, 517
Spermatozoa, in urine, 504
Sperm-oil, 458
*Spiræa ulmaria*, 420
**Spirit** of French wine, 426
methylated, 434
myrcia, 423
of nitrous ether, 349, 430
of nitrous ether, adulterated, 435
proof, 423
pyroxylic, 434
rectified, 423
of turpentine, 466
of wine, 423
of wine, impurities in, 622
wood-, 434
Spirits, 507
analysis of, 610
*Spiritus ætheris*, 430
*ætheris nitrosi*, 430
*ætheris nitrosi*, impurities in, 621
*ammoniæ*, 93
*ammoniæ aromaticus*, 93
*ammoniæ aromaticus*, impurities in, 621
*ammoniæ fœtidus*, 93
*anisi*, 462
*cajuputi*, 462
*cinnamomi*, 462
*chloroformi*, impurities in, 621
*frumenti*, 423
*juniperi*, 462
*lavandulæ*, 462
*menthæ piperitæ*, 462
*myrciæ*, 423
*myristicæ*, 462

*Spiritus*—
*rectificatus*, 423, 557
*rectificatus*, impurities in, 622
*rosmarini*, 462
*tenuior*, 423
*tenuior*, impurities in, 622
*vini gallici*, 426
Spodumene, 227
Spogel seeds, 406
Spongy platinum, 248
Spontaneous combustion, 158
Spotted cranesbill, 357
Spruce fir, 470
Spurge laurel, 470
*Squalus carcharius*, 460
Squill, 412
Standard gold, 245
**Standard solution** of hyposulphite of sodium, 564
solution of iodine, 558
solution of nitrate of silver, 556
solution of oxalic acid, 546
solution of red chromate of potassium, 561
solution of soda, 553
solution of sulphuric acid, 551
Stannate of sodium, 243
Stannates, 242
Stannic acid, 242
anhydride, 242
chloride, 242
oxide, 242
sulphide, 243
anhydrous, 244
Stannous chloride, solid, 242
hydrate, 243
oxide, 243
sulphide, 243
*Stannum*, 34
Star-anise oil, 463
Starch, 401
blue, 401
quantitative estimation of, 608
-sugar, 411
Starches, microscopy of, 401
Stas's process for poisonous alkaloids, 490
Stavesacre, 396
Steam-bath, 569
Stearic acid, 458, 460
Stearine, 458
Steatite, 478
Steel, 141
wine, 155
Stick liquorice, 418
*Stibium*, 32
Still, 127
Stoddart's test for quinine, 386
Stone-coal, 241
red, 476
Storax, 473
Stout, 423
*Stramonii folia*, 393
*semina*, 393
Strasburg turpentine, 466
Strawberry, sugar in, 410
Stream-tin, 241
Strontianite, 228
**Strontium**, 228
analytical reactions of, 228
carbonate of, 228
derivation of word, 33
flame, 229
nitrate of, 228
sulphate of, 229
Structure of flame, 23
*Strychnia*, 389
*Strychniæ sulphas*, 390
Strychnia or strychnine, 389
or strychnine, impurities in, 622
analytical reactions of, 390 488
in organic mixtures, detection of, 488
*Strychnos Ignatius*, 389
*Strychnos nux vomica*, 389
Styracin, 473
*Styrax benzoin*, 473
*præparatus*, 474
Styrol, 474
Styrone, 473
Subacetate of copper, 192
of lead, 211
Subcarbonate of iron, 144
Subchloride of mercury, 202
Sublimation, 92, 201
Sublimed sulphur, 362
Subnitrate of bismuth, 251
Substances readily deoxidized, quantitative estimation of, 563
readily oxidized, quantitative estimation of, 558
Substitution products, 437
Succi, 507

Succinate of potassium, 354
Succinic acid, 354
*Succinum*, 354
*Succus limonum*, 325
Sucrose, 410
Suet, 458
prepared, 458
Sugar, 409
amount in various fruits, 410
brown, 410
-candy, 410
cane-, 410
detection of, in urine, 494
grape, 410
inverted, 409
lump, 410
-maple, 410
milk-, 411
moist, 410
of gelatine, 457
of lead, 211
quantitative estimation of, 608
tests for, 409
**Sulphate** of aluminium, 138
aluminium and ammonium, 137
ammonium, 90
ammonium and iron, 138
barium, 102
bismuth, 253
cadmium, 249
calcium, 105, 114
chromium, 238
cinchonia, 388
cobalt, 234
copper, 192
copper, anhydrous, 192
cupr-diammon-diammonium, 206
indigo, 290
iron, 142
iron, solution of, 143
lead, 214
magnesium, 117, 576
manganese, 233
mercury, 199
potassium, 73, 287
quinine, 384
sodium, 87, 268
strontium, 228
zinc, 130
Sulphates, 308
analytical reactions of, 311
Sulphates—
quantitative estimation of, 590
Sulphethylic acid, 428
**Sulphide** of allyl, 447
ammonium, 95
antimony, 179, 182
arsenicum, 167, 176
arsenicum, native, 167
barium, 102
bismuth, 254
cadmium, 249
calcium, 303
cobalt, 234
copper, 192
iron, 145
lead, 214
lead, native, 209
manganese, 232
mercury, 207
mercury, native, 194
nickel, 236
potassium, 68
silver, 219
silver, native, 216
tin, 243
zinc, 135
zinc, native, 130
Sulphides, 301
analytical reactions of, 304
native, 167
quantitative estimation of, 589
Sulphindigotic acid, 290
Sulphindylic acid, 290
**Sulphite** of barium, 307
calcium, 306
magnesium, 305
potassium, 305
silver, 306
sodium, 305
Sulphites, 305
analytical reactions of, 307
quantitative estimation of, 590
Sulphocarbolates, 445
Sulphocarbolic acid, 445
Sulphocyanate of acrinyl, 447
allyl, 447
butyl, 434
iron, 161
Sulphocyanates, 354
Sulphocyanic acid, 354
Sulphocyanides, 355
Sulphocyanogen, 355

Sulphophenates, 445
Sulphophenic acid, 445
Sulphosalicylic acid, 446
Sulphostannates, 244
Sulphovinic acid, 428
**Sulphur**, 30
adulteration of, 304
allotropy of, 302
analytical reactions of, 304
arsenic in, 176
bromide of, 305
chloride of, 305
derivation of word, 31
estimation of, 589
flowers of, 302
hypochloride of, 305
iodide of, 274, 305
liver of, 68
milk of, 304
oxyacids, 345
plastic, 302
precipitated, 303
roll, 302
sublimed, 302
*Sulphur lotum*, 302
*præcipitatum*, 303
*præcipitatum*, impurities in, 622
*sublimatum*, 302
*sublimatum*, impurities in, 622
Sulphurated antimony, 182
potash, 68
Sulphurets, *vide* Sulphides.
Sulphuretted hydrogen, 303
Sulphuric acid, 308
acid and sulphuric anhydride in dilute acid of different densities, Tables showing the percentage of, 624
acid, antidotes to, 312
acid, aromatic, 310
acid, dilute, 310
acid, fuming, 310
acid, Nordhausen, 310
acid in organic mixtures, detection of, 486
acid, purification of, 310
standard solution of, 551
acid, volumetric estimation of, 554
anhydride, 310
*Sulphuris iodidum*, 274
*iodidum*, impurities in, 622
Sulphurous acid, 30, 305
Sulphurous—
acid, volumetric estimation of, 559
anhydride, 305
Sulphydrate of ammonium, solution of, 95
Sulphydric acid, 96, 303
Sumatra camphor, 467
Sumbul, 472
Superphosphate of lime, 327
Supporters of combustion, 22
*Suppositoria acidi tannici*, 356
*morphiæ*, 382
*plumbi composita*, 211
Suppositories, 507
Surface, unit, 521
Surgery, 14
Sweetbread, 455
Sweet spirit of nitre, 349, 430
spirit of nitre, adulterated, 435
Sylvic acid, 469
Symbol, function of, 41, 52
Symbols of elements, 30, 41
illustration of chemical action by, 46
Sympathetic inks, 235
Synaptase, 414
Synthesis, 60
Syphon, 110
Syrup of iodide of iron, 30
Syrups, 507
specific gravities of, 532
*Syrupi*, impurities in, 622
*Syrupus aurantii*, 463
*aurantii floris*, 463
*ferri iodidi*, 30
*ferri phosphatis*, 145
*fuscus*, 412

*Tabaci folia*, 397
Tables, various (*see* Appendix).
Talc, 137
*Tamarindus*, 324
impurities in, 622
Tannic acid, 355
Tanning, 356
Tantalum, 636
Tapioca, 401
starch (fig.), 403
*Taraxaci radix*, 400
Taraxacin, 400
Tartar, meaning of, 319
Tartar, cream of, 72, 79, 319
emetic, 181, 319

Tartar—
emetic, estimation of antimony in, 579
Tartarated antimony, 181
Tartaric acid, 318, 319
saturating power of, 321, 623
solution of, 320
*Tartarus boraxatus*, 334
**Tartrate** of ammonium, 98
antimony and potassium, 181, 319
calcium, 322
potassium, acid, 73, 79
potassium, neutral, 73, 320
potassium and sodium, 85, 320
silver, 322
sodium, 79
**Tartrates**, 73, 318
analytical reactions of, 322
volumetric estimation of, 554
Taurine, 457
Taurocholates, 457
Teal oil, 460
Teleni fly, 468
Tellurium, 303, 638
Temperature, correction of vol. of gas for, 536
measurement of, 513
*Terebinthina*, 466
*canadensis*, 466
Terpenes, 461
Terra di sienna, 476
*Terra japonica*, 357
*Testa præparata*, 110
Test-papers, 95
-tube, 16
Tetramines, 380
Tetrathionic acid, 345
Tetryl, 433
Thalleiochin, 386
Thallium, 636
Thebaia, 382
Theia, 398
Theine, 398
Thenard's blue, 477
Theobroma oil, 458
Theobromate of glyceryl, 458
Therapeutics, definition and derivation of, 14
*Theriaca*, 412
Thermolysis, 539
Thermometer, 513
Thermometer—
Celsius's, 514
Centigrade, 514
Fahrenheit's, 514
Reaumur's, 514
Thermometric scales, conversion of degrees of, 514
Thionic acids, 345
Thorinum, 636
Thorium, 636
Thorn-apple, 393
*Thus americanum*, 471
Thyme, oil of, 466
Thymene, 466
Thymol, 463, 466
*Thymus vulgaris*, 466
Tiglic acid, 459
**Tin**, 241
amalgam, 241
analytical reactions of, 243
antidotes to, 244
block, 241
chloride of, 242
derivation of word, 34
dropped or grain, 241
foil, 241
granulated, 241
oxide of, 242
perchloride of, 242
plate, 241
prepare-liquor, 243
-stone, 241
tacks, 241
-white cobalt, 234
Tincal, 333
*Tinctura ferri acetatis*, 149
*ferri perchloridi*, 148
*iodi*, 274
*iodinii*, 274
*iodinii composita*, 274
*quiniæ*, 385
*quiniæ ammoniata*, 385
*Tincturæ*, 148
Tinctures, 148, 507
Tinnevelly senna, 415
*Tinospora cordifolia*, 400
Titanium, 636
Tobacco, 397
*Toddaliæ radix*, 400
Tolen, 473
Tolu, balsam of, 473
Tolnol, 443
Toxicology, 483

Tragacanth, 114
*Tragacantha*, 114
Treacle, 412
Triads, 122
Triamines, 380
Triangle, wire, 99
Tribasic acids, 265
Tribasylous radicals, 265
Triethylamine, 379
Triethylia, 379
Trinitro-carbolic acid, 444
Trinitrocellulin, 408
Triphane, 227
Tripoli, 352
Trithionic acid, 345
*Triticum*, starch of (fig.), 403
Trityl, 433
Tritylia, 379
Trivalence, 56
Trivalent radicals, 56, 122
*Trochisci acidi tannici*, 356
*bismuthi*, 252
*ferri redacti*, 158
*morphiæ*, 382
*morphiæ et ipecacuanhæ*, 382
*potassæ chloratis*, 294
*sodæ bicarbonatis*, 85
Tropate of atropia, 393
Tropic acid, 393
Tropine, 393
Tube-funnels, 97
Tubes for collecting gases, 18
glass, *vide* Glass tubes.
Tungsten, 636
Turgite, 151
Turkey corn, 396
Turmeric, 475
oil, 466
paper, 95, 334
Turnbull's blue, 341, 477
Turpentine, 466
American, 466
Bordeaux, 466
Canadian, 466
Chian, 466
French, 466
rectified oil of, 466
spirit of, 466
Strasburg, 466
Venice, 466
Turpeth mineral, 200
Turps, 466
*Tylophora asthmatica*, 396
Type-metal, 179, 209
Types, chemical, 377
Typical formulæ, 433

*Ulmi cortex*, 357
*fulvæ cortex*, 357
Ultimate analysis, 596
Ultramarine blue, 477
green, 477
Umbelliferone, 472
Umber, 477
*Uncaria gambier*, 357
*Unguentum aconitiæ*, 393
*antimonii tartarati*, 182
*atropiæ*, 393
*cadmii iodidi*, 249
*cerussæ*, 210
*hydrargyri*, 195
*hydrargyri ammoniati*, 206
*hydrargyri iodidi rubri*, 198
*hydrargyri nitratis*, 199
*hydrargyri oxidi rubri*, 203
*hydrargyri subchloridi*, 202
*iodi*, 274
*iodinii compositum*, 274
*plumbi acetatis*, 211
*plumbi carbonatis*, 210
*plumbi iodidi*, 212
*plumbi subacetatis compositum*, 211
*sulphuris iodidi*, 274
*veratriæ*, 399
Units of capacity, 521
surface, 521
weight, 521
Univalence, 56
Univalent radicals, 55, 122
Uranium, 636
Urari, 391
Urate of lithium, 228
Urates, 358
*Urceola elastica*, 474
Urea, 337, 495
artificial, 337, 495
Uric acid, 358, 498
Urinary calculi, 504
calculi, examination of, 504
deposits or sediments, plates of, *vide* 498 *et seq.*
sediments, 496
sediments, microscopical examination of, 499
Urine, 493
diabetic, 410, 494, 608
estimation of urea in, 495

Urine—
morbid, examination of, 493
Urinometer, 533
*Uvæ*, 412
*ursi folia*, 357

Valerene, 467
Valerian oil, 467
*Valerianæ radix*, 467
Valerianate of amyl, 441
quinia, 485
sodium, 360
zinc, 134, 360
Valerianates, 359
Valerianic acid, 359, 460
Valerol, 467
Vanadates, 332
Vanadinite, 332
Vanadium, 332, 636
relationship to nitrogen, phosphorus, and arsenicum, 332
Vanilla, 473
Vanillic acid, 361
Vanilline, 361
*Vapor acidi hydrocianici*, 282
*chlori*, 29
*coniæ*, 396
*iodi*, 273
Vapor-density, 536
*Variolaria*, 477
Vegetable albumen, 453
and animal life, relation of, 19
casein, 401, 453
crocus, 475
fibrin, 401, 453
gelatine, 453
green, 477
jelly, 405
oil, 456
rouge, 476
substances, 378 *et seq.*
Venetian red, 151
Venice turpentine, 466
Veratralbia, 397
*Veratri viridi radix*, 397
*Veratria*, 399
impurities in, 622
Veratria or Veratrine, 399
*Veratrum album*, 397
*viride*, 397
Verdigris, 192
Vermilion, 207
*Veronica virginica*, 400

Vinegar, 297
estimation of mineral acids in, 590
impurities in, 613
of cantharides, 298
of squill, 298
*Vinum antimoniale*, 182
*aurantii*, 423
*ferri*, 155
*ferri citratis*, 155
*portense*, 423
*quiniæ*, 385
*xericum*, 423
Vitriol, blue, 143
green, 143
oil of, 310
white, 143
Volatile oils, *vide* Oils.
Volatility of salts of ammonium, 98
Volatilization, 99
Volcanic ammonia, 90
Volume, combination by, 52
of gas, corrections of, 526
molecular, 54
**Volumetric** analysis, 543
estimation of acetate of lead, 546
estimation of acetic acid, 554
estimation of acids, 552
estimation of alkalies, 544
estimation of alkaline carbonates, 550
estimation of ammonia solutions, 546
estimation of arseniate of iron, 562
estimation of arseniate of sodium, 556
estimation of arsenic and arsenical solutions, 559
estimation of borax, 546
estimation of bromides, 556
estimation of bromide of potassium, 556
estimation of chlorides, 585
estimation of chlorinated lime, 565
estimation of chlorinated soda, 566
estimation of chlorine, 564
estimation of citrate of potassium, 550
estimates of citrates, 551

Volumetric—
estimation of cyanides, 556
estimation of hydrochloric acid, 554
estimation of hydrocyanic acid, 556
estimation of hyposulphite of sodium, 560
estimation of iodide of potassium, 557
estimation of magnetic oxide of iron, 562
estimation of nitric acid, 554
estimation of official compounds, 546, 554, 556, 559, 562, 564
estimation of phosphate of iron, 562
estimation of potash, 550
estimation of saccharated carbonate of iron, 562
estimation of soda, 547
estimation of sugar, 608
estimation of sulphides, 589
estimation of sulphites, 596
estimation of sulphuric acid, 554
estimation of sulphurous acid, 559
estimation of tartrates, 550
solutions, 546, 553, 556, 558, 564
Vulcanite, 474
Vulcanized India-rubber, 474

Wahoo bark, 400
Washing-bottles, 109, 569
precipitates, 109, 570
Warmth of animals, how kept up, 19
Water, 127
aërated, 85
ammonia in, potable, 548
aspirator, 309
-bath, 569
boiling-point of, 515
chalybeate, 140
composition of, 22
crystallization, 84
quantitative estimation of, 613
cubic inches of, in a gallon, 537
distilled, 127
Water—
evaporation of, 70
formation of, expressed by symbols, 43
hardness of, 315
hemlock, 464
lime-, 107
nitrites in, 349
of crystallization, 84
-oven, 569
oxygenated, 103
purification of, 126, 315
softness of, 315
type, 377
weight of 1 cubic inch of, 537
weight of minim, drachm, ounce, pint, and gallon, 526
Wax, 458
Weighing-tubes, 568
Weight, 518
estimation of, 518
molecular, 54
of air, 537
of hydrogen, 538
of water, 529
specific, 529
Weights, atomic, 51
and measures of the British Pharmacopœia of 1867, 525
and measures of the metric decimal system, 522 *et seq.*
and measures of the U. S. Pharmacopœia of 1870, 519
balance, 519
relation of metrical to the weights of the U. S. Pharmacopœia, 525
relative, 52
Weld, 465
Welding, 248
Wheat starch (fig.), 403
Wheaten flour, 401
Whey, 411, 451
Whiskey, 423
White arsenic, 167
indigo, 291
lead, 210
pepper, 397
pigments, 478
precipitate, fusible, 206
infusible, 206
resin, 469
vitriol, 143
wax, 458, 517

Whiting, 110
Whortleberry, sugar in, 412
Wild indigo, 400
Willow bark, 346, 420
Wine, 423, 610
  antimonial, 182
  iron, 155
  orange, 423
  quinine, 385
  sherry, 423
  steel, 155
Winter-green, oil of, 442
Wire-gauze tray, 28
  triangle, 99
Witherite, 102
Wood-charcoal, 112
  creasote, 444
  naphtha, 434
  oil, 471
  spirit, 434
  tar, 471
Woody nightshade, 398
Wormseed, 467
Wourali, 391
Wourara, 391
Wrought iron, 141

Xanthin, 506
*Xanthorrhiza apiifolia*, 394
*Xanthoxylon fraxineum*, 394
Xylol, 443

Yard, 526
Yeast, 422
Yelk of egg, 450
Yellow chromate of potassium, 237
  coloring-matters, 475
  dock, 337
  ochre, 475
  oxide of mercury, 203
  parilla, 394
  prussiate of potass., 279, 339
  root, 394
  sienna, 475
  wax, 458, 517
  wood, 475
Yolk of egg, 450
Yttrium, 636

Zaffre, 234
Zanaloin, 446
Zanzibar aloes, 446
Zea, starch of (fig.), 403
**Zinc**, 130
  acetate of, 133
  analytical reactions of, 135
  antidotes to, 136
  carbonate of, 130, 132
  chloride of, 131
  derivation of word, 32
  detection of, in presence of aluminium and iron, 162
  -ethyl, 432
  ferrocyanide of, 135
  granulated, 20
  hydrate of, 135
  in organic mixtures, detection of, 486
  oxide of, 133
  oxide of, Hubbuck's, 134
  quantitative estimation of, 576
  sulphate of, 130
  sulphide of, 135
  sulphide of, native, 130
  valerianate of, 134, 360
  white, 133
*Zinci acetas*, 133
  *acetas*, impurities in, 622
  *carbonas*, 130, 132
  *carbonas*, impurities in, 622
  *chloridi, liquor*, 132
  *chloridum*, 131
  *chloridum*, impurities in, 622
  *oxidum*, 133
  *oxidum*, impurities in, 622
  *sulphas*, 130
  *sulphas*, impurities in, 622
  *valerianas*, 134, 360
  *valerianas*, impurities in, 622
*Zincum*, 130
  *granulatum*, 20
*Zingiber*, 467
Zirconium, 636

# EXTRACTS FROM CRITIQUES.

## Great Britain.

THE LANCET.—"We thoroughly believe that the course of study prescribed by this book will teach the student the principles of chemistry and will give him a special knowledge concerning the applications of that science to pharmacy..... We commend it to the notice of every medical as well as pharmaceutical student."

THE CHEMICAL NEWS.—"For all the numerous class of students who are preparing for the medical or for the pharmaceutical profession, we know of no work in the language which can be compared with the one before us."

THE PHARMACEUTICAL JOURNAL.—"It is, indeed, a most useful manual. It is a necessary companion to every student following the system of practical instruction at the Society's laboratory; it is also the best possible guide to the pharmaceutical pupil who cannot indulge in laboratory tuition."

THE MEDICAL TIMES AND GAZETTE.—"It is a good work for beginning the study of theoretical chemistry. It is the same as regards practical chemistry.....and an admirable companion to the British Pharmacopœia. It is rare to find so many qualities combined, and quite curious to note how much valuable information finds a mutual interdependence."

THE BRITISH MEDICAL JOURNAL.—"Dr. Attfield's volume, just published, is rather a new book than a second edition of his previous work, and more nearly realizes our ideal than any book we have before seen on the subject."

THE CHEMIST AND DRUGGIST.—"Professor Attfield's 'Introduction to Pharmaceutical Chemistry' has developed into a handsome volume of nearly 650 pages, and is now a complete text-book of chemistry, adapted for use as a reading-book, a lecture-companion, and a laboratory guide."

THE MEDICAL PRESS AND CIRCULAR —"We believe that this Manual has been already adopted as the class-book by many of the professors in the public schools throughout the United Kingdom.....In pharmaceutical chemistry applied to the Pharmacopœia, we know of no rival. It is, therefore, particularly suited to the medical student."

DUBLIN QUARTERLY JOURNAL OF MEDICAL SCIENCE.—"Dr. Attfield's book contains more information of a practical character than is to be found in most works of its class. It is just the kind of chemical manual which is most suited to the wants of the student of medicine.....We cordally recommend it."

NATURE.—"We have derived much satisfaction from the perusal of Dr. Attfield's work; it is eminently practical in its character, and it is written with a just appreciation of the small amount of time for the study of chemistry at the disposal of the student in medicine and pharmacy."

## France.

MONITEUR SCIENTIFIQUE.—"Dans un in-12 de plus de 600 pages l'auteur a magnifiquement développé une chimie minérale et organique en l'appliquant à deux sciences qui lui sont solidaires, la médecine et la pharmacie.....Nous pouvons dire qu'il est indispensable aux chimistes par la variété des matières qu'il renferme et par le choix judicieux que l'auteur a apporté de tout ce qui se rattache à la chimie.....Nous espérons qu'une traduction française viendra augmenter le nombre des lecteurs de son ouvrage"

JOURNAL DE PHARMACIE ET DE CHIMIE.—"Le plan adopté par le savant professeur a un caractère original, qui lui assure une place distinguée parmi les ouvrages destinés à répandre la connaissance de la chimie.....Nous avons en France nombre de bons ouvrages destinés à faire connaître la chimie, mais nous sommes persuadés que, traduit en notre langue, le livre de M. Attfield pourrait rivaliser avec eux et ne compterait pas moins des lecteurs."

## Germany.

NEUES REPERTORIUM FUR PHARMACIE.—"In der That zeichnet sich dieses Werk, wie wir uns überzeugt haben, durch seinen reichen Inhalt und hohe Originalität ebensosehr wie durch leichte Verständlichkeit und durch die geschickte Art, womit darin Theorie und Praxis verbunden sind, aus."

ARCHIV DER PHARMACIE.—"Es enthält, auf 48 Druckbogen zusammengedrängt, Alles, was Studirende der Medecin und Pharmacie nöthig wissen mussen.....Die Art unde Weise, wie der Verfasser den grossen Stoff behandelt, ist ebenso instructiv, als originell."

[*Continued on next page.*]

# EXTRACTS FROM CRITIQUES (*continued*).

## America.

PHILADELPHIA MEDICAL TIMES.—"Certainly the bugbears of those medical students whose opportunities for the study of chemical philosophy have been limited, are dispelled in a way which merits their sincere thanks: while a great deal of the confusion which seems to exist as regards atoms and molecules, quantivalence, and equivalent weights will be removed from their minds, and withal so gradually, that the wonder will be that it ever existed. In its mode of teaching, the book fills a place hitherto vacant."

CINCINNATI MEDICAL REPERTORY.—"This is decidedly the best work on chemistry for medical students with which we have ever met; and when we assert this, we wish to be understood as meaning precisely what we say. There are many excellent works upon chemistry, but very few, if any of them, have been prepared with special reference to the wants of the medical student; but this one has, and it is that which constitutes its excellence. We would expect that it would be used in our medical colleges as a text-book, to the exclusion of all others."

LEAVENWORTH MEDICAL HERALD AND JOURNAL OF PHARMACY.—"It is one of the most instructive volumes which has come under our notice, and specially adapted to the wants of every pharmaceutist. We are particularly pleased with the practical manner in which medicinal preparations are treated with reference to the United States Pharmacopœia. Our readers will find it of special interest."

AMERICAN JOURNAL OF PHARMACY.—"A more careful examination of this work has fully confirmed the opinion which we expressed in our last number. It is a valuable guide to the medical and pharmaceutical student who, by practical experiments, desires to gain a thorough knowledge of chemistry. If we consider the large scope and the small size of the work, it is astonishing what an immense number of facts the author succeeded in embracing in this space. We heartily recommend this work to the pharmacist and physician."

NEW YORK MEDICAL RECORD.—"The author has taken pains to revise his work with special reference to the wants of American students, introducing the materia medica of the United States Pharmacopœia, and making sundry other additions, which altogether render it one of the most desirable text-books on the subject of chemistry that the student can obtain."

VIRGINIA CLINICAL RECORD.—"The work is a most valuable one. Such a manual of the chemistry of the Pharmacopœia, has been long needed, and will prove of great assistance not only to students, but to the practising pharmacist and physician."

BOSTON JOURNAL OF CHEMISTRY.—"The work holds a high rank in England, and as a handbook for the medical student and the pharmacist we know of nothing better in English."

CHICAGO PHARMACIST.—"Wherever practical chemistry is taught (and it should be in every college), Attfield's manual will be found not simply useful, but indispensable. We regard it as by far the best work of the kind that has come under our notice, and as such we heartily commend it to our readers."

NEW YORK MEDICAL GAZETTE.—"Most of our readers have probably found, from time to time, difficulty in hunting out from general treatises on chemistry the practical points of information which they have needed for direct medical use. To all such we take pleasure in commending Dr. Attfield's excellent manual, which is designed for their especial benefit."

THE AMERICAN JOURNAL OF THE MEDICAL SCIENCES.—"Dr. Attfield's work will prove a useful assistant, not only to those who desire to study the subject in its practical bearings, but also to the general student in chemistry, to the apothecary, and to the physician."

## Canada.

CANADIAN MEDICAL JOURNAL.—"It in every way fulfils the intention of the author. We can strongly recommend it as a most complete manual of chemistry, alike useful to the physician and pharmaceutist."

CANADIAN PHARMACEUTICAL JOURNAL.—"We would simply recommend this work as the best with which we are acquainted, either as accompanying a course of lectures on pharmacopœial chemistry, or as a guide to the self-taught student. To the pharmacist it will prove a very useful counter companion; indeed, we may say that the shop library is incomplete without it."

---

**HENRY C. LEA'S SON & CO.—Philadelphia.**

# CATALOGUE OF BOOKS

PUBLISHED BY

# HENRY C. LEA'S SON & CO.

(LATE HENRY C. LEA.)

---

The books in the annexed list will be sent by mail, post-paid, to any Post Office in the United States, on receipt of the printed prices. No risks of the mail, however, are assumed, either on money or books. Gentlemen will, therefore, in most cases, find it more convenient to deal with the nearest bookseller.

*In response to a large number of inquiries for a finer binding than is usually placed on medical books, we now finish many of our standard publications in half Russia, using in the manufacture none but the best materials. To foster the growing taste, the prices have been fixed at so small an advance over the cost of leather binding as to bring it within the reach of all to possess a library attractive to the eye as well as to the mind.*

Detailed catalogues furnished or sent free by mail on application.

HENRY C. LEA'S SON & CO.,
Nos. 706 and 708 Sansom Street, Philadelphia.

---

## PERIODICALS.

## THE MEDICAL NEWS,

### A WEEKLY JOURNAL OF MEDICAL SCIENCE,

*FIVE DOLLARS ($5) per annum, in advance,*

*Is published every Saturday, commencing January 7, 1882, and contains from 28 to 32 large double-columned quarto pages, the contributors to which include the most able writers of the day.*

### THE AMERICAN JOURNAL OF THE MEDICAL SCIENCES,

*FIVE DOLLARS ($5) per annum, in advance,*

EDITED BY I. MINIS HAYS, A.M., M.D.,

*Is published quarterly, on the 1st of January, April, July and October. Each number contains over 300 large octavo pages.*

*COMMUTATION RATES.*

THE MEDICAL NEWS
THE AMERICAN JOURNAL OF MEDICAL SCIENCES } Nine dollars per annum, in advance.

---

ALLEN (HARRISON). A SYSTEM OF HUMAN ANATOMY. WITH AN INTRODUCTORY CHAPTER ON HISTOLOGY, by E. O. Shakespeare, M.D To be completed in one large and handsome quarto volume of about 550 double-columned pages, with 380 engravings on stone on 110 plates, and 250 woodcuts in the text. In six sections, each in a portfolio. Section I. (Histology) and Section II. (Bones and Joints) just ready. Each $3 50.

*Sold by subscription only.*

AN AMERICAN SYSTEM OF GYNÆCOLOGY. In Treatises by Various Authors. Edited by Horatio R. Bigelow, M.D. (*In active preparation.*)

ASHHURST (JOHN, Jr.) THE PRINCIPLES AND PRACTICE OF SURGERY. FOR THE USE OF STUDENTS AND PRACTITIONERS. Third and revised edition. In one large octavo volume of over 1000 pages, containing about 550 woodcuts. (*In press*)

ASHTON (T. J.) ON THE DISEASES, INJURIES AND MALFORMATIONS OF THE RECTUM AND ANUS. With remarks on Habitual Constipation. Second American from the fourth London edition, with illustrations. One 8vo. vol. of 287 pp. Cloth, $3 25.

ASHWELL (SAMUEL). A PRACTICAL TREATISE ON THE DISEASES OF WOMEN. Third American from the third London edition. In one octavo volume of 520 pages. Cloth, $3 50.

ATTFIELD (JOHN). CHEMISTRY; GENERAL, MEDICAL AND PHARMACEUTICAL. Eighth edition, revised by the Author. In One 12mo. vol. of 701 pages, with 87 illus. Clo., $2 50; leather, $3 00.

BAIRD (ROBERT). IMPRESSIONS AND EXPERIENCES OF THE WEST INDIES. 1 vol. royal 12mo. Cloth, 75 cents.

BARLOW (GEORGE H.) A MANUAL OF THE PRACTICE OF MEDICINE. In one 8vo. volume of 603 pages. Cloth, $2 50.

BARNES (FANCOURT). A MANUAL OF MIDWIFERY FOR MIDWIVES. In one royal 12mo. vol., with numerous illustrations. Cloth, $1 25. (*Just issued.*)

BARNES (ROBERT). A PRACTICAL TREATISE ON THE DISEASES OF WOMEN. Third Am. from third Eng. edition. In one handsome 8vo. vol. of about 800 pp., with about 200 illustrations. (*Preparing.*)

BARTHOLOW (ROBERTS). A PRACTICAL TREATISE ON ELECTRICITY AS APPLIED TO THE PRACTICE OF MEDICINE. Second edition. In one 8vo. vol. of about 300 pages. (*Preparing.*)

BASHAM (W. R.) RENAL DISEASES; A CLINICAL GUIDE TO THEIR DIAGNOSIS AND TREATMENT. In one 12mo. volume of 304 pages, with illustrations. Cloth, $2 00.

BELLAMY'S MANUAL OF SURGICAL ANATOMY. In one royal 12mo. volume of 300 pages, with 50 illustrations. Cloth, $2 25.

BLOXAM (C. L.) CHEMISTRY, INORGANIC AND ORGANIC. With Experiments. In one handsome octavo volume of 700 pages, with 295 illustrations. Cloth, $4 00; leather, $5 00.

BRISTOWE (JOHN SYER). A TREATISE OF THE PRACTICE OF MEDICINE. Second American edition, revised by the Author. Edited with additions by James H. Hutchinson, M.D. In one handsome 8vo. volume of 1085 pages. Cloth, $5 00; leather, $6 00; very handsome half Russia, raised bands, $6 50. (*Just issued.*)

BROWNE (EDGAR A.) HOW TO USE THE OPHTHALMOSCOPE. Elementary instruction in Ophthalmoscopy for the Use of Students. In one small 12mo. volume of 116 pages, with 35 illust. Cloth, $1.

BROWNE (LENNOX). THE THROAT AND ITS DISEASES. New edition. In one handsome imperial 8vo. volume, with one hundred illustrations in color and fifty woodcuts. (*Preparing.*)

BRYANT (THOMAS). THE PRACTICE OF SURGERY. Third American from the third English edition. Edited, with additions, by J. B. Roberts, M.D. In one imperial octavo volume of 1009 pages, with 735 illustrations. Cloth, $6 50; leather, $7 50; very handsome half Russia, raised bands, $8. (*Just issued.*)

BUMSTEAD (F. J.) THE PATHOLOGY AND TREATMENT OF VENEREAL DISEASES. Fourth edition, revised and enlarged, with the co-operation of R. W. Taylor, M.D. 1 8vo. vol. of 843 pages, with 138 illustrations. Cloth, $4 75; leather, $5 75; very handsome half Russia, raised bands, $6 25. (*Just issued.*)

——AND CULLERIER'S ATLAS OF VENEREAL. See "CULLERIER."

BURNETT (CHARLES H.) THE EAR: ITS ANATOMY, PHYSIOLOGY AND DISEASES. A Practical Treatise for the Use of Students and Practitioners. In one handsome octavo volume of 619 pages, with 87 illustrations. Cloth $4 50; leather, $5 50; very handsome half Russia, raised bands, $6. (*Lately issued.*)

CAMPBELL'S LIVES OF LORDS KENYON, ELLENBOROUGH AND TENTERDEN. Being the third volume of "Campbell's Lives of the Chief Justices of England." In one crown octavo vol. Cloth, $2.

CARPENTER (WM. B.) PRINCIPLES OF HUMAN PHYSIOLOGY. A new American, from the eighth English edition. In one large 8vo. volume of 1083 pages, with 373 illustrations. Cloth, $5 50; leather, raised bands, $6 50; very handsome half Russia, raised bands, $7. (*Lately issued.*)

——— PRIZE ESSAY ON THE USE OF ALCOHOLIC LIQUORS IN HEALTH AND DISEASE. New Edition, with a Preface by D. F. Condie, M.D. One 12mo. volume of 178 pages. Cloth, 60 cents.

CARTER (R. BRUDENELL). A PRACTICAL TREATISE ON DISEASES OF THE EYE. With additions and test-types, by John Green, M.D. In one 8vo. vol. of 500 pages, with 124 illustrations.

CENTURY OF AMERICAN MEDICINE.—A HISTORY OF MEDICINE IN AMERICA, 1776–1876. In one 12mo. vol. of 366 pages. Cloth, $2 25.

CHADWICK (JAMES R.) A MANUAL OF THE DISEASES PECULIAR TO WOMEN. In one 12mo. vol., with illust. (*Preparing.*)

CHAMBERS (T. K.) A MANUAL OF DIET IN HEALTH AND DISEASE. In one handsome 8vo. vol. of 302 pages. Cloth, $2 75.

CHURCHILL (FLEETWOOD). ESSAYS ON THE PUERPERAL FEVER. In one octavo volume of 464 pages. Cloth, $2 50.

CLASSEN'S QUANTITATIVE ANALYSIS. Translated by Edgar F. Smith, Ph.D. In one 12mo. volume of 324 pp., with 36 illustrations. Cloth, $2 00.

CLELAND (JOHN). A DIRECTORY FOR THE DISSECTION OF THE HUMAN BODY. In one 12mo. vol. of 178 pp. Cloth, $1 25.

CLOWES (FRANK). AN ELEMENTARY TREATISE ON PRACTICAL CHEMISTRY AND QUALITATIVE INORGANIC ANALYSIS. From the Third Eng. Ed. In one 12mo. vol. Cloth, $2 50.

COLEMAN (ALFRED). A MANUAL OF DENTAL SURGERY AND PATHOLOGY. With Notes and Additions to adapt it to American Practice. By Thos. C. Stellwagen, M.A., M.D., D.D.S. In one handsome 8vo. vol. of 412 pp., with 331 illus. Cloth, $3 25. (*Just ready.*)

CONDIE (D. FRANCIS). A PRACTICAL TREATISE ON THE DISEASES OF CHILDREN. Sixth edition, revised and enlarged. In one large 8vo. vol. of 800 pages. Cloth, $5 25; leather, $6 25.

COOPER (B.B.) LECTURES ON THE PRINCIPLES AND PRACTICE OF SURGERY. In one large 8vo. vol. of 767 pages. Cloth, $2 00.

CORNIL (V.) SYPHILIS: ITS MORBID ANATOMY, DIAGNOSIS AND TREATMENT. Translated, with notes and additions, by J. Henry C. Simes, M.D., and J. William White, M.D. In one 8vo. volume of 461 pages, with 84 illust. Cloth, $3 75. (*Just ready.*)

CORNIL (V.), AND RANVIER (L.) MANUAL OF PATHOLOGICAL HISTOLOGY. Translated, with Notes and Additions, by E. O. Shakespeare, M.D., and J. Henry C. Simes, M.D. In one octavo volume of 800 pages, with 360 illustrations. Cloth, $5 50; leather, $6 50; very handsome half Russia, raised bands, $7. (*Just issued.*)

COURTY (A.) A PRACTICAL TREATISE ON THE DISEASES OF THE UTERUS, THE OVARIES AND THE FALLOPIAN TUBES. Translated by Agnes McLaren, M.D., M.R.C.P.I. With an introduction by J. Matthews Duncan, M.D., LL.D., F.R.S.E. (*Preparing.*)

CULLERIER (A.) AN ATLAS OF VENEREAL DISEASES. Translated and edited by FREEMAN J. BUMSTEAD, M.D. A large imperial quarto volume, with 26 plates containing about 150 figures, beautifully colored, many of them the size of life. In one vol., strongly bound in cloth, $17. In five parts, paper covers, $3 per part.

DALTON (J. C.) A TREATISE ON HUMAN PHYSIOLOGY. Seventh edition, thoroughly revised, and greatly improved. In one very handsome 8vo. vol. of 722 pages, with 252 illustrations. Cloth, $5; leather, $6; very handsome half Russia, $6 50. (*Just ready.*)

DANA (JAMES D.) THE STRUCTURE AND CLASSIFICATION OF ZOOPHYTES. With illust. on wood. In one imp. 4to. vol. Clo., $4.

DAVIS (F. H.) LECTURES ON CLINICAL MEDICINE. Second ed., revised and enlarged. In one 12mo. vol. of 287 pp. Cloth, $1 75.

DE LA BECHE'S GEOLOGICAL OBSERVER. In one large 8vo. vol. of 700 pages, with 300 illustrations. Cloth, $4.

DON QUIXOTE DE LA MANCHA. Illustrated edition. In two handsome crown 8vo. vols. Cloth, $2 50; half morocco, $3 70.

DRUITT (ROBERT). THE PRINCIPLES AND PRACTICE OF MODERN SURGERY. A revised American, from the eighth London edition. In one octavo volume of 687 pages, with 432 wood engravings. Cloth, $4; leather, $5.

DUNCAN (J. MATTHEWS). CLINICAL LECTURES ON THE DISEASES OF WOMEN. Delivered in St. Bartholomew's Hospital. In one octavo volume of 175 pages. Cloth, $1 50. (*Just issued.*)

DUNGLISON (ROBLEY). MEDICAL LEXICON; A Dictionary of Medical Science. Containing a concise explanation of the various subjects and terms of Anatomy, Physiology, Pathology, Hygiene, Therapeutics, Pharmacology, Pharmacy, Surgery, Obstetrics, Medical Jurisprudence and Dentistry; notices of Climate and of Mineral Waters; Formulæ for Officinal, Empirical and Dietetic Preparations; with the accentuation and Etymology of the Terms, and the French and other Synonymes. New edition. In one very large royal 8vo. vol. Cloth, $6 50; leather, $7 50; half Russia, raised bands, $8.

EDIS (ARTHUR W.). DISEASES OF WOMEN. A Manual for Students and Practitioners. In one handsome 8vo. vol., of 576 pp., with 148 illustrations. Cloth, $3; leather, $4. (*Just ready.*)

ELLIS (GEORGE VINER). DEMONSTRATIONS IN ANATOMY. Being a Guide to the Knowledge of the Human Body by Dissection. From the eighth and revised English edition. In one very handsome octavo volume of 716 pages, illustrated by 249 engravings on wood. Cloth, $4 25; leather, $5 25. (*Lately issued.*)

EMMET (THOMAS ADDIS). THE PRINCIPLES AND PRACTICE OF GYNÆCOLOGY, for the use of Students and Practitioners. Second edition, enlarged and revised. In one large 8vo. vol. of 879 pp., with 133 original illustrations. Cloth, $5; leather, $6; very handsome half Russia, raised bands, $6 50. (*Lately issued.*)

ERICHSEN (JOHN E.) THE SCIENCE AND ART OF SURGERY. A new American, from the seventh enlarged and revised London edition. Specially revised by the Author. In two large octavo volumes of 1944 pages, with 863 illustrations. Cloth, $8 50; leather, $10 50; very handsome half Russia, raised bands, $11 50.

FARQUHARSON (ROBERT). A GUIDE TO THERAPEUTICS. Third American edition, specially revised by the Author. Edited, with additions, embracing the U. S. Pharmacopœia, by Frank Woodbury, M.D. In one royal 12mo. volume. (*In Press.*)

FENWICK (SAMUEL). THE STUDENTS' GUIDE TO MEDICAL DIAGNOSIS. From the third revised and enlarged London edition. In one royal 12mo. volume of 328 pages. Cloth, $2 25.

FINLAYSON (JAMES). A CLINICAL MANUAL FOR THE STUDY OF MEDICAL CASES. In one handsome 8vo. vol. of 546 pages, with 85 woodcuts. Cloth, $2 63. (*Lately issued.*)

FLINT (AUSTIN). A TREATISE ON THE PRINCIPLES AND PRACTICE OF MEDICINE. Fifth edition, thoroughly revised and largely rewritten. In one large 8vo. vol. of 1150 pages Cloth, $5 50; leather, $6 50; very handsome half Russia, $7. (*Just issued.*)

——— A MANUAL OF PERCUSSION AND AUSCULTATION; of the Physical Diagnosis of Diseases of the Lungs and Heart, and of Thoracic Aneurism. Second edition, revised and enlarged. In one handsome royal 12mo. volume. Cloth, $1 63

——— A PRACTICAL TREATISE ON THE DIAGNOSIS AND TREATMENT OF DISEASES OF THE HEART. Second edition, enlarged. In one octavo volume of 550 pages. Cloth, $4 00.

——— A PRACTICAL TREATISE ON THE PHYSICAL EXPLORATION OF THE CHEST, AND THE DIAGNOSIS OF DISEASES AFFECTING THE RESPIRATORY ORGANS. Second and revised edition. In one octavo volume of 591 pages. Cloth, $4 50.

——— CLINICAL MEDICINE. A SYSTEMATIC TREATISE ON THE DIAGNOSIS AND TREATMENT OF DISEASE. Designed for Students and Practitioners of Medicine. In one handsome octavo volume of 799 pages. Cloth, $4 50; leather, $5 50; very handsome half Russia, raised bands, $6 00 (*Lately issued.*)

——— MEDICAL ESSAYS. In one 12mo. vol., pp. 210. Cloth, $1 38.

——— ON PHTHISIS: ITS MORBID ANATOMY, ETIOLOGY, ETC., in a series of Clinical Lectures. A new work. In one handsome 8vo. volume. Cloth, $3 50.

FOSTER (MICHAEL). TEXT-BOOK OF PHYSIOLOGY. Second Am. from the last London Edition, with extensive notes and additions by E. T. Reichert, M D. In one large 12mo. vol. of 999 pages, with 259 illustrations. Cloth, $3 25; leather, $3 75. (*Just Ready.*)

——— TEXT-BOOK OF PHYSIOLOGY. Latest English Edition. In one handsome 12mo. vol. of 804 pp., with 72 illus. Cloth, $3 00.

FOTHERGILL'S PRACTITIONER'S HANDBOOK OF TREATMENT. Second edition, revised and enlarged. In one very neat octavo volume of about 650 pp. Cloth, $4 00; very handsome half Russia, $5 50. (*Just issued.*)

FOWNES (GEORGE). A MANUAL OF ELEMENTARY CHEMISTRY. A new American, from the twelfth English edition. In one royal 12mo. vol. of 1031 pages, with 177 illustrations, and one colored plate. Cloth, $2 75; leather, $3 25. (*Just issued.*)

FOX (TILBURY). EPITOME OF SKIN DISEASES, with Formulæ for Students and Practitioners. Third Am. edition, specially revised by the author. In one small 12mo. vol. (*Preparing*)

FULLER (HENRY). ON DISEASES OF THE LUNGS AND AIR PASSAGES. Their Pathology, Physical Diagnosis, Symptoms and Treatment. From 2d Eng. ed. In 1 8vo. vol., pp. 475. Cloth, $3 50.

GALLOWAY (ROBERT). A MANUAL OF QUALITATIVE ANALYSIS. From the sixth London edition. (*Preparing.*)

GIBSON'S INSTITUTES AND PRACTICE OF SURGERY. In two octavo volumes of about 1000 pages. Leather, $6 50.

GLUGE (GOTTLIEB). ATLAS OF PATHOLOGICAL HISTOLOGY. Translated by Joseph Leidy, M.D., Professor of Anatomy in the University of Pennsylvania, &c. In one imperial quarto vol., with 320 copperplate figures, plain and colored. Cloth, $4.

GRAY (HENRY). ANATOMY, DESCRIPTIVE AND SURGICAL. A new American, from the eighth and enlarged London edition. To which is added Holden's "Landmarks, Medical and Surgical," with additions by W. W. Keen, M.D. In one imperial octavo volume of 993 pages, with 523 large and elaborate engravings on wood. Cloth, $6; leather, $7; very handsome half Russia, raised bands, $7 50.

GREEN (T. HENRY). AN INTRODUCTION TO PATHOLOGY AND MORBID ANATOMY. Fourth American, from the fifth London edition. In one handsome octavo volume of 351 pages, with 138 illustrations. Cloth, $2 25. (*Now ready.*)

GREENE (WILLIAM H.) A MANUAL OF MEDICAL CHEMISTRY. For the Use of Students. Based upon Bowman's Medical Chemistry. In one royal 12mo. volume of 310 pages, with 74 illustrations. Cloth, $1 75. (*Just issued.*)

GRIFFITH (ROBERT E.) A UNIVERSAL FORMULARY, CONTAINING THE METHODS OF PREPARING AND ADMINISTERING OFFICINAL AND OTHER MEDICINES. Third and enlarged edition. Edited by John M. Maisch, Phar.D. In one large 8vo. vol. of 800 pages, double columns. Cloth, $4 50; leather, $5 50.

GROSS (SAMUEL D.) A PRACTICAL TREATISE ON THE DISEASES, Injuries and Malformations of the Urinary Bladder, the Prostate Gland and the Urethra. Third edition, thoroughly revised and much condensed, by Samuel W. Gross, M.D. In one octavo volume of 574 pages, with 170 illus. Cloth, $4 50. (*Lately issued.*)

——— A SYSTEM OF SURGERY, PATHOLOGICAL, DIAGNOSTIC, THERAPEUTIC AND OPERATIVE. Fifth edition, revised and improved. In two imperial 8vo. vols. containing 2276 pages, with 1402 illustrations. Strongly bound in leather, raised bands, $15; very handsome half Russia, raised bands, $16.

——— A PRACTICAL TREATISE ON FOREIGN BODIES IN THE AIR PASSAGES. In one 8vo. vol. of 468 pages. Cloth, $2 75.

GROSS (SAMUEL W.) A PRACTICAL TREATISE ON IMPOTENCE, STERILITY, AND ALLIED DISORDERS OF THE MALE SEXUAL ORGANS. In one handsome octavo volume of 166 pages, with 16 illustrations. Cloth, $1 50. (*Now ready.*)

GUSSEROW (A.) A TREATISE ON UTERINE TUMORS. Revised by the Author, and translated by Edmund C. Wendt, M.D., with additions by the translator and Nathan Bozeman, M.D. In one 8vo. volume of about 400 pages, with many illustrations. (*Preparing.*)

HABERSHON (S. O.) ON THE DISEASES OF THE ABDOMEN, AND OTHER PARTS OF THE ALIMENTARY CANAL. Second American, from the third English edition. In one handsome 8vo. volume of 554 pages, with illus. Cloth, $3.50. (*Just issued.*)

HALL (MRS. M.) LIVES OF THE QUEENS OF ENGLAND BEFORE THE NORMAN CONQUEST. In one handsome 8vo. vol. Cloth, $2 25; crimson cloth, $2 50; half morocco, $3.

HAMILTON (ALLAN McLANE) NERVOUS DISEASES, THEIR DESCRIPTION AND TREATMENT. Second and revised edition. In one handsome octavo volume of 598 pages, with 72 illustrations. Cloth, $4. (*Just ready.*)

HAMILTON (FRANK H.) A PRACTICAL TREATISE ON FRACTURES AND DISLOCATIONS. Sixth edition, thoroughly revised. In one handsome 8vo. vol. of 909 pages, with 352 illus. Cloth, $5 50; leather, $6 50; very handsome half Russia, $7. (*Just issued.*)

**HARTSHORNE (HENRY).** ESSENTIALS OF THE PRINCIPLES AND PRACTICE OF MEDICINE. Fifth edition. In one 12mo. vol. 669 pp. with 144 illus. Cl., $2 75; half bound, $3. (*Just ready.*)

——— A HANDBOOK OF ANATOMY AND PHYSIOLOGY. In one royal 12mo. volume of 310 pages, with 220 illust. Cloth, $1 75.

——— CONSPECTUS OF THE MEDICAL SCIENCES. Comprising Manuals of Anatomy, Physiology, Chemistry, Materia Medica, Practice of Medicine, Surgery and Obstetrics. Second edition. In one royal 12mo. volume of 1028 pages, with 477 illustrations. Cloth, $4 25; leather, $5 00.

**HEATH (CHRISTOPHER).** PRACTICAL ANATOMY; A MANUAL OF DISSECTIONS. With additions, by W. W. Keen, M. D.

**HILL (BERKELEY).** SYPHILIS AND LOCAL CONTAGIOUS DISORDERS In one 8vo. volume of 479 pages. Cloth, $3 25.

**HILLIER (THOMAS).** HAND-BOOK OF SKIN DISEASES. 2d ed. In one royal 12mo. volume of 353 pp., with two plates. Cloth, $2 25.

**HOBLYN (RICHARD D.).** A DICTIONARY OF THE TERMS USED IN MEDICINE AND THE COLLATERAL SCIENCES. In one 12mo. vol. of 520 double-columned pp. Cloth, $1 50; leather, $2.

**HODGE (HUGH L.)** ON DISEASES PECULIAR TO WOMEN, INCLUDING DISPLACEMENTS OF THE UTERUS. Second and revised edition. In one 8vo. volume of 519 pages. Cloth, $4 50.

——THE PRINCIPLES AND PRACTICE OF OBSTETRICS. In one large 4to. vol. of 542 double-columned pages, illustrated with large lithographic plates containing 159 figures from original photographs, and 110 woodcuts. Strongly bound in cloth, $14.

**HOFFMANN (FREDERICK) AND POWER (FREDERICK B.)** A MANUAL OF CHEMICAL ANALYSIS, as Applied to the Examination of Medicinal Chemicals and their Preparations. Third edition, entirely rewritten and much enlarged. In one handsome octavo volume, fully illustrated. (*In press.*)

**HOLDEN (LUTHER).** LANDMARKS, MEDICAL AND SURGICAL. From the third English edition. With additions, by W. W. Keen, M.D. In one royal 12mo. vol. of 148 pp. Cloth, $1. (*Just ready.*)

**HOLLAND (SIR HENRY).** MEDICAL NOTES AND REFLECTIONS. From 3d English ed. In one 8vo. vol. of about 500 pp. Cloth, $3 50.

**HOLMES (TIMOTHY.)** A SYSTEM OF SURGERY. From the last English edition, with notes and additions by various American authors. Edited by John H. Packard, M.D. In three very handsome 8vo. vols. containing 3137 double-columned pp., with 979 illus. and 13 lithographic plates. Cloth, $18; leather, $21; very handsome half Russia, raised bands, $22 50. (*Just ready.*) *For sale by subscription only.*

**HOLMES (TIMOTHY).** SURGERY, ITS PRINCIPLES AND PRACTICE. In one handsome octavo volume of 968 pages, with 411 illustrations. Cloth, $6; leather, raised bands, $7; very handsome half Russia, raised bands, $7 50.

**HORNER (WILLIAM E.)** SPECIAL ANATOMY AND HISTOLOGY. Eighth edition, revised and modified. In two large 8vo. vols. of 1007 pages, containing 320 woodcuts. Cloth, $6.

**HUDSON (A.)** LECTURES ON THE STUDY OF FEVER. In one octavo volume of 308 pages. Cloth, $2 50.

**HUGHES.** SCRIPTURE GEOGRAPHY AND HISTORY, with 12 colored maps. In one 12mo. volume. Cloth, $1.

**HYDE (JAMES NEVINS).** A PRACTICAL TREATISE ON DISEASES OF THE SKIN. In one handsome octavo volume of about 600 pages, with many illustrations. (*In press.*)

JONES (C. HANDFIELD). CLINICAL OBSERVATIONS ON FUNCTIONAL NERVOUS DISORDERS. Second American edition. In one octavo volume of 340 pages. Cloth, $3 25.

KEATING (JOHN M.) THE CARE OF INFANTS. In one small 12mo. volume of 118 pages. Cloth, $1. (*Now ready.*)

KING (A F. A.). A MANUAL OF OBSTETRICS. In one very handsome 12mo. volume of 321 pages, with 58 illustrations. Cloth. $2 00. (*Just ready.*)

KNAPP (F.) TECHNOLOGY, OR CHEMISTRY APPLIED TO THE ARTS AND TO MANUFACTURES; with American additions, by Prof. Walter R. Johnson. In two 8vo. vols., with 500 ill. Cloth. $6.

LA ROCHE (R.) YELLOW FEVER. In two 8vo. vols. of 1468 pages. Cloth, $7.

——— PNEUMONIA. In one 8vo. vol. of 490 pages. Cloth, $3.

LAURENCE (J. Z.) AND MOON (ROBERT C.) A HANDY-BOOK OF OPHTHALMIC SURGERY. Second edition, revised by Mr Laurence. In one 8vo. vol. pp. 227, with 66 illus. Cloth, $2 75

LAWSON (GEORGE). INJURIES OF THE EYE, ORBIT AND EYELIDS. From the last English edition. In one handsome octavo volume of 404 pages, with 92 illustrations. Cloth, $3 50.

LEA (HENRY C.) SUPERSTITION AND FORCE; ESSAYS ON THE WAGER OF LAW, THE WAGER OF BATTLE, THE ORDEAL AND TORTURE. Third edition, thoroughly revised and greatly enlarged. In one handsome royal 12mo. vol. pp. 552. Cloth, $2 50.

——— STUDIES IN CHURCH HISTORY. The Rise of the Temporal Power—Benefit of Clergy—Excommunication. In one handsome 12mo. vol. of 519 pp.

——— AN HISTORICAL SKETCH OF SACERDOTAL CELIBACY IN THE CHRISTIAN CHURCH. In one handsome octavo volume of 602 pages. Cloth, $2 50.

LEE (HENRY) ON SYPHILIS. In one 8vo. vol. pp. 246. Cloth, $2 25.

LEHMANN (C. G.) PHYSIOLOGICAL CHEMISTRY. Translated by George F. Day, M. D. In two large 8vo. volumes containing 1195 pages, with plates, and 168 woodcuts. Cloth, $6.

——— A MANUAL OF CHEMICAL PHYSIOLOGY. In one very handsome 8vo. vol. of 327 pages, with 41 woodcuts. Cloth, $2 25.

LEISHMAN (WILLIAM). A SYSTEM OF MIDWIFERY. Including the Diseases of Pregnancy and the Puerperal State. Third American, from the third English edition. With additions, by J. S. Parry, M.D. In one very handsome octavo volume of 740 pages, with 205 illustrations. Cloth, $4 50; leather, $5 50; very handsome half Russia, $6. (*Lately issued.*)

LUDLOW (J. L.) A MANUAL OF EXAMINATIONS UPON ANATOMY, PHYSIOLOGY, SURGERY, PRACTICE OF MEDICINE, OBSTETRICS, MATERIA MEDICA, CHEMISTRY, PHARMACY AND THERAPEUTICS. To which is added a Medical Formulary. Third edition. In one royal 12mo. volume of 816 pages, with 370 woodcuts. Cloth, $3 25; leather, $3 75.

LYNCH (W. F.) A NARRATIVE OF THE UNITED STATES EXPEDITION TO THE DEAD SEA AND RIVER JORDAN. In one large octavo vol., with 28 beautiful plates and two maps. Cloth, $3.

——— Same Work, condensed edition. One vol. royal 12mo. Cloth, $1.

LYONS (ROBERT D.) A TREATISE ON FEVER. In one octavo volume of 362 pages. Cloth, $2 25.

MACFARLAND'S TURKEY AND ITS DESTINY. In 2 vols. royal 12mo. Cloth, $2.

MAISCH (JOHN M.) A MANUAL OF ORGANIC MATERIA MEDICA. In one handsome 12mo. volume of 451 pages, with 194 beautiful illustrations. Cloth, $2 75. (*Just ready.*)

MARSH (MRS.) A HISTORY OF THE PROTESTANT REFORMATION IN FRANCE. In 2 vols. royal 12mo. Cloth, $2.

MEIGS (CHAS. D.) ON THE NATURE, SIGNS AND TREATMENT OF CHILDBED FEVER. In one 8vo. vol. of 346 pages. Cloth, $2.

MILLER (JAMES). PRINCIPLES OF SURGERY. Fourth American, from the third Edinburgh edition. In one large octavo volume of 688 pages, with 240 illustrations. Cloth, $3 75.

——— THE PRACTICE OF SURGERY. Fourth American, from the last Edinburgh edition. In one large octavo volume of 682 pages, with 364 illustrations. Cloth, $3 75.

MITCHELL (S. WEIR). LECTURES ON NERVOUS DISEASES, ESPECIALLY IN WOMEN. Second edition. (*Preparing.*)

MONTGOMERY (W. F.) AN EXPOSITION OF THE SIGNS AND SYMPTOMS OF PREGNANCY. From the second English edition. In one handsome 8vo. vol. of 568 pages, with illus. Cloth, $3 75.

MORRIS (MALCOLM). SKIN DISEASES: Including their Definitions, Symptoms, Diagnosis, Prognosis, Morbid Anatomy and Treatment. A Manual for Students and Practitioners In one 12mo. vol. of 316 pages, with illus. Cloth, $1 75. (*Just issued.*)

MULLER (J.) PRINCIPLES OF PHYSICS AND METEOROLOGY. In one large 8vo. vol. of 623 pages, with 538 cuts. Cloth, $4 50.

NEILL (JOHN) AND SMITH (FRANCIS G.) A COMPENDIUM OF THE VARIOUS BRANCHES OF MEDICAL SCIENCE. In one handsome 12mo. volume of 974 pages, with 374 woodcuts. Cloth, $4; leather, raised bands, $4 75.

NETTLESHIP'S MANUAL OF OPHTHALMIC MEDICINE In one royal 12mo. vol. of 377 pages, with 89 illustrations. Cloth, $2. (*Just issued.*)

PAGET'S HUNGARY AND TRANSYLVANIA. In two royal 12mo. volumes. Cloth, $2.

PARRISH (EDWARD). A TREATISE ON PHARMACY. With many Formulæ and Prescriptions. Fourth edition. Enlarged and thoroughly revised by Thomas S. Wiegand. In one handsome octavo volume of 985 pages, with 280 illustrations. Cloth, $5 50; leather, $6 50; very handsome half Russia, raised bands, $7.

PARVIN (THEOPHILUS). A TREATISE ON MIDWIFERY. In one handsome 8vo. vol. of about 550 pp., with many illus. (*Preparing.*)

PAVY (F. W.) A TREATISE ON THE FUNCTION OF DIGESTION, ITS DISORDERS AND THEIR TREATMENT. From the second London edition. In one octavo volume of 238 pages. Cloth, $2.

PIRRIE (WILLIAM). THE PRINCIPLES AND PRACTICE OF SURGERY. In one handsome octavo volume of 780 pages, with 316 illustrations. Cloth, $3 75.

PLAYFAIR (W. S.) A TREATISE ON THE SCIENCE AND PRACTICE OF MIDWIFERY. Third American edition, revised by the author. Edited, with Additions, by R. P. Harris, M.D. In one handsome octavo volume of 659 pages, with 183 woodcuts, and two plates. Cloth, $4; leather, $5; very handsome half Russia, raised bands, $5 50. (*Just issued.*)

POLITZER (ADAM). A TEXT-BOOK OF AURAL THERAPEUTICS for Practising Surgeons and Students. Translated with notes and additions. In one handsome octavo volume of about 600 pages, with about 200 illustrations. (*Preparing.*)

PULSZKY'S MEMOIRS OF AN HUNGARIAN LADY. In one royal 12mo. vol. Cloth, $1.

RAMSBOTHAM (FRANCIS H.) THE PRINCIPLES AND PRACTICE OF OBSTETRIC MEDICINE AND SURGERY. In one imperial octavo volume of 640 pages, with 64 plates, besides numerous woodcuts in the text. Strongly bound in leather, $7.

REMSEN (IRA). THE PRINCIPLES OF CHEMISTRY. In one handsome 12mo. volume of 232 pages. Cloth, $1 50.

REYNOLDS (J. RUSSELL). A SYSTEM OF MEDICINE, with Notes and Additions, by HENRY HARTSHORNE, M.D. In three large 8vo. vols., containing 3056 closely printed double-columned pages, with 317 illus. Per vol., cloth, $5; leather, $6; very handsome half Russia, $6 50. (*Now ready.*) *For sale by subscription only.*

RICHARDSON (BENJAMIN W.) PREVENTIVE MEDICINE. In one octavo volume of about 500 pages. (*Preparing.*)

ROBERTS (JOHN B.) THE PRINCIPLES AND PRACTICE OF SURGERY. In one octavo volume of about 500 pages, fully illustrated. (*Preparing.*)

ROBERTS (WILLIAM). A PRACTICAL TREATISE ON URINARY AND RENAL DISEASES. Fourth American, from the fourth London edition. With numerous illustrations and a colored plate. In one very handsome 8vo. vol. of over 600 pages. (*Preparing.*)

SARGENT (F. W.) ON BANDAGING AND OTHER OPERATIONS OF MINOR SURGERY. New edition, with an additional chapter on Military Surgery. In one handsome royal 12mo. volume of 383 pages, with 187 woodcuts. Cloth, $1 75.

SCHÄFER (EDWARD ALBERT). A COURSE OF PRACTICAL HISTOLOGY: A Manual of the Microscope for Medical Students. In one handsome octavo volume, with many illustrations. Cloth, $2.

SCHMITZ AND ZUMPT'S CLASSICAL SERIES. In royal 18mo.

ADVANCED LATIN EXERCISES, WITH SELECTIONS FOR READING. Revised. Cloth, price 60 cents; half bound, 70 cents.

C. C. SALLUSTII DE BELLO CATILINARIO ET JUGURTHINO. With notes, map, &c. Price in cloth, 60 cents; half bound, 70 cents.

CORNELII NEPOTIS LIBER DE EXCELLENTIBUS DUCIBUS EXTERARUM GENTIUM, CUM VITIS CATONIS ET ATTICI. With notes, &c. Price in cloth, 60 cents; half bound, 70 cts.

P. VIRGILII MARONIS CARMINA OMNIA. Price in cloth, 85 cents; half bound, $1.

Q. CURTII RUFI DE GESTIS ALEXANDRI MAGNI LIBRI VIII. With notes, map, &c. Price in cloth, 80 cents; half bound, 90 cents.

SCHOEDLER (FREDERICK) AND MEDLOCK (HENRY). WONDERS OF NATURE. An elementary introduction to the Sciences of Physics, Astronomy, Chemistry, Mineralogy, Geology, Botany, Zoology and Physiology. Translated from the German by H. Medlock. In one 8vo. vol., with 679 illustrations. Cloth, $3.

SEILER (CARL). HANDBOOK OF DIAGNOSIS AND TREATMENT OF DISEASES OF THE THROAT AND NASAL CAVITIES. Second edition. (*Preparing.*)

SHARPEY (WILLIAM) AND QUAIN (JONES AND RICHARD). HUMAN ANATOMY. With notes and additions by Jos. Leidy, M.D., Prof. of Anatomy in the University of Pennsylvania. In two large 8vo. vols. of about 1300 pages, with 511 illustrations. Cloth, $6.

SKEY (FREDERIC C.) OPERATIVE SURGERY. In one 8vo. vol. of over 650 pages, with 81 woodcuts. Cloth, $3 25.

SLADE (D. D.) DIPHTHERIA; ITS NATURE AND TREATMENT. Second edition. In one royal 12mo. vol. pp. 158. Cloth, $1 25.

SMALL BOOKS ON GREAT SUBJECTS. In 3 vols. Cloth, $1 50.

SMITH (EDWARD). CONSUMPTION; ITS EARLY AND REMEDIABLE STAGES. In one 8vo. vol. of 253 pp. Cloth, $2 25.

SMITH (EUSTACE). ON THE WASTING DISEASES OF CHILDREN. Second American edition, enlarged. In one 8vo. vol. Cloth, $2 50.

SMITH (HENRY H.) AND HORNER (WILLIAM E.) ANATOMICAL ATLAS. Illustrative of the structure of the Human Body. In one large imperial 8vo. vol., with about 650 beautiful figures. Clo., $4 50.

SMITH (J. LEWIS). A TREATISE ON THE DISEASES OF INFANCY AND CHILDHOOD. Fifth edition, revised and enlarged. In one large 8vo. volume of 836 pages, with illustrations. Cloth, $4 50; leather, $5 50; very handsome half Russia, raised bands, $6. (*Just ready.*)

STILLÉ (ALFRED). THERAPEUTICS AND MATERIA MEDICA. Fourth revised edition. In two handsome octavo volumes of 1936 pages. Cloth, $10; leather, $12; very handsome half Russia, $13.

STILLÉ (ALFRED) AND MAISCH (JOHN M.) THE NATIONAL DISPENSATORY: Embracing the Chemistry, Botany, Materia Medica, Pharmacy, Pharmacodynamics and Therapeutics of the Pharmacopœias of the United States and Great Britain. For the Use of Physicians and Pharmaceutists. Second edition, revised and enlarged. In one handsome octavo volume of 1692 pages, with 239 illustrations. Cloth, $6 75; leather, $7 50; very handsome half Russia, raised bands, $8 25. (*Just issued.*)

STIMSON (LEWIS A.) A PRACTICAL TREATISE ON FRACTURES. In one handsome octavo volume of about 600 pages, fully illustrated. (*In press.*)

——— A MANUAL OF OPERATIVE SURGERY. In one royal 12mo. volume of 477 pages, with 332 illustrations. Cloth, $2 50.

STOKES (W.) LECTURES ON FEVER. In one 8vo. vol. Cloth, $2.

STRICKLAND (AGNES). LIVES OF THE QUEENS OF HENRY VIII. AND OF HIS MOTHER. In one crown octavo vol. Extra cloth, $1; black cloth, 90 cents.

——— MEMOIRS OF ELIZABETH, SECOND QUEEN REGNANT OF ENGLAND AND IRELAND. In one crown octavo vol. Extra cloth, $1 40; black cloth, $1 30.

STURGES (OCTAVIUS). AN INTRODUCTION TO THE STUDY OF CLINICAL MEDICINE. In one 12mo. vol. Cloth, $1 25.

TANNER (THOMAS HAWKES). A MANUAL OF CLINICAL MEDICINE AND PHYSICAL DIAGNOSIS. Third American from the second revised English edition. Edited by Tilbury Fox, M. D. In one handsome 12mo. volume of 362 pp., with illus. Cloth, $1 50.

——— ON THE SIGNS AND DISEASES OF PREGNANCY. From the second English edition. In one 8vo. volume of 490 pages, with four colored plates and numerous woodcuts. Cloth, $4 25.

TAYLOR (ALFRED S.) MEDICAL JURISPRUDENCE. Eighth American from tenth English edition, revised by the author. Edited by John J. Reese, M.D. In one large octavo volume of 937 pages, with 70 illustrations. Cloth, $5; leather, $6; very handsome half Russia, raised bands, $6 50. (*Now ready.*)

——— ON POISONS IN RELATION TO MEDICINE AND MEDICAL JURISPRUDENCE. Third American from the third London edition. In one octavo volume of 788 pages, with 104 illustrations. Cloth, $5 50; leather, $6 50. (*Just issued.*)